



Cochrane
Library

Cochrane Database of Systematic Reviews

Chemotherapy for advanced gastric cancer (Review)

Wagner AD, Syn NLX, Moehler M, Grothe W, Yong WP, Tai BC, Ho J, Unverzagt S

Wagner AD, Syn NLX, Moehler M, Grothe W, Yong WP, Tai BC, Ho J, Unverzagt S.
Chemotherapy for advanced gastric cancer.

Cochrane Database of Systematic Reviews 2017, Issue 8. Art. No.: CD004064.

DOI: [10.1002/14651858.CD004064.pub4](https://doi.org/10.1002/14651858.CD004064.pub4).

www.cochranelibrary.com

Chemotherapy for advanced gastric cancer (Review)

Copyright © 2017 The Cochrane Collaboration. Published by John Wiley & Sons, Ltd.

WILEY

TABLE OF CONTENTS

ABSTRACT	1
PLAIN LANGUAGE SUMMARY	2
SUMMARY OF FINDINGS	4
BACKGROUND	22
OBJECTIVES	23
METHODS	23
RESULTS	26
Figure 1.	27
Figure 2.	31
Figure 3.	33
DISCUSSION	40
AUTHORS' CONCLUSIONS	44
ACKNOWLEDGEMENTS	46
REFERENCES	47
CHARACTERISTICS OF STUDIES	68
DATA AND ANALYSES	149
Analysis 1.1. Comparison 1 Chemotherapy versus best supportive care, Outcome 1 Overall survival.	149
Analysis 1.2. Comparison 1 Chemotherapy versus best supportive care, Outcome 2 Time to progression.	150
Analysis 2.1. Comparison 2 Combination versus single-agent chemotherapy, Outcome 1 Overall survival.	150
Analysis 2.2. Comparison 2 Combination versus single-agent chemotherapy, Outcome 2 Tumour response.	151
Analysis 2.3. Comparison 2 Combination versus single-agent chemotherapy, Outcome 3 Time to progression.	152
Analysis 2.4. Comparison 2 Combination versus single-agent chemotherapy, Outcome 4 Treatment-related death.	152
Analysis 3.1. Comparison 3 5-FU/cisplatin/anthracycline combinations versus 5-FU/cisplatin combinations (without anthracyclines), Outcome 1 Overall survival.	153
Analysis 3.2. Comparison 3 5-FU/cisplatin/anthracycline combinations versus 5-FU/cisplatin combinations (without anthracyclines), Outcome 2 Tumour response.	153
Analysis 3.3. Comparison 3 5-FU/cisplatin/anthracycline combinations versus 5-FU/cisplatin combinations (without anthracyclines), Outcome 3 Time to progression.	153
Analysis 4.1. Comparison 4 5-FU/cisplatin/anthracycline combinations versus 5-FU/anthracycline combinations (without cisplatin), Outcome 1 Overall survival.	154
Analysis 5.1. Comparison 5 Chemotherapy with irinotecan versus non-irinotecan-containing regimes, Outcome 1 Overall survival.	155
Analysis 5.2. Comparison 5 Chemotherapy with irinotecan versus non-irinotecan-containing regimes, Outcome 2 Tumour response.	156
Analysis 5.3. Comparison 5 Chemotherapy with irinotecan versus non-irinotecan-containing regimes, Outcome 3 Progression-free survival.	156
Analysis 5.4. Comparison 5 Chemotherapy with irinotecan versus non-irinotecan-containing regimes, Outcome 4 Treatment-related death.	157
Analysis 5.5. Comparison 5 Chemotherapy with irinotecan versus non-irinotecan-containing regimes, Outcome 5 Treatment discontinuation due to toxicity.	158
Analysis 6.1. Comparison 6 Chemotherapy with docetaxel versus non-docetaxel-containing regimes, Outcome 1 Overall survival.	159
Analysis 6.2. Comparison 6 Chemotherapy with docetaxel versus non-docetaxel-containing regimes, Outcome 2 Tumour response.	159
Analysis 6.3. Comparison 6 Chemotherapy with docetaxel versus non-docetaxel-containing regimes, Outcome 3 Time to progression.	160
Analysis 6.4. Comparison 6 Chemotherapy with docetaxel versus non-docetaxel-containing regimes, Outcome 4 Progression-free survival.	161
Analysis 6.5. Comparison 6 Chemotherapy with docetaxel versus non-docetaxel-containing regimes, Outcome 5 Treatment-related death.	161
Analysis 6.6. Comparison 6 Chemotherapy with docetaxel versus non-docetaxel-containing regimes, Outcome 6 Treatment discontinuation due to toxicity.	162
Analysis 7.1. Comparison 7 Chemotherapy with capecitabine versus 5-FU-containing regimes, Outcome 1 Overall Survival.	162

Analysis 7.2. Comparison 7 Chemotherapy with capecitabine versus 5-FU-containing regimes, Outcome 2 Tumour response. ..	163
Analysis 7.3. Comparison 7 Chemotherapy with capecitabine versus 5-FU-containing regimes, Outcome 3 Time to progression.	163
Analysis 7.4. Comparison 7 Chemotherapy with capecitabine versus 5-FU-containing regimes, Outcome 4 Progression-free survival.	163
Analysis 7.5. Comparison 7 Chemotherapy with capecitabine versus 5-FU-containing regimes, Outcome 5 Treatment-related death.	164
Analysis 7.6. Comparison 7 Chemotherapy with capecitabine versus 5-FU-containing regimes, Outcome 6 Treatment discontinuation due to toxicity.	164
Analysis 8.1. Comparison 8 Chemotherapy with oxaliplatin versus the same regime including cisplatin, Outcome 1 Overall Survival.	164
Analysis 8.2. Comparison 8 Chemotherapy with oxaliplatin versus the same regime including cisplatin, Outcome 2 Tumour response.	165
Analysis 8.3. Comparison 8 Chemotherapy with oxaliplatin versus the same regime including cisplatin, Outcome 3 Progression-free survival.	165
Analysis 8.4. Comparison 8 Chemotherapy with oxaliplatin versus the same regime including cisplatin, Outcome 4 Treatment-related death.	166
Analysis 8.5. Comparison 8 Chemotherapy with oxaliplatin versus the same regime including cisplatin, Outcome 5 Treatment discontinuation due to toxicity.	166
Analysis 9.1. Comparison 9 Taxane-platinum-fluoropyrimidine combinations versus taxane-platinum (without fluoropyrimidine), Outcome 1 Overall survival.	167
Analysis 9.2. Comparison 9 Taxane-platinum-fluoropyrimidine combinations versus taxane-platinum (without fluoropyrimidine), Outcome 2 Tumour response.	167
Analysis 9.3. Comparison 9 Taxane-platinum-fluoropyrimidine combinations versus taxane-platinum (without fluoropyrimidine), Outcome 3 Progression-free survival.	167
Analysis 9.4. Comparison 9 Taxane-platinum-fluoropyrimidine combinations versus taxane-platinum (without fluoropyrimidine), Outcome 4 Treatment-related death.	168
Analysis 9.5. Comparison 9 Taxane-platinum-fluoropyrimidine combinations versus taxane-platinum (without fluoropyrimidine), Outcome 5 Treatment discontinuation due to toxicity.	168
Analysis 10.1. Comparison 10 S-1 versus 5-FU-containing regimes, Outcome 1 Overall Survival.	169
Analysis 10.2. Comparison 10 S-1 versus 5-FU-containing regimes, Outcome 2 Tumour response.	169
Analysis 10.3. Comparison 10 S-1 versus 5-FU-containing regimes, Outcome 3 Progression-free survival.	169
Analysis 10.4. Comparison 10 S-1 versus 5-FU-containing regimes, Outcome 4 Time-to treatment failure.	170
Analysis 10.5. Comparison 10 S-1 versus 5-FU-containing regimes, Outcome 5 Treatment-related deaths.	170
Analysis 10.6. Comparison 10 S-1 versus 5-FU-containing regimes, Outcome 6 Treatment discontinuation due to toxicity.	170
APPENDICES	171
WHAT'S NEW	175
HISTORY	175
CONTRIBUTIONS OF AUTHORS	175
DECLARATIONS OF INTEREST	175
SOURCES OF SUPPORT	176
DIFFERENCES BETWEEN PROTOCOL AND REVIEW	176
INDEX TERMS	177

[Intervention Review]

Chemotherapy for advanced gastric cancer

Anna Dorothea Wagner¹, Nicholas LX Syn², Markus Moehler³, Wilfried Grothe⁴, Wei Peng Yong², Bee-Choo Tai⁵, Jingshan Ho², Susanne Unverzagt⁶

¹Department of Oncology, Lausanne University Hospitals and Clinics, Lausanne, Switzerland. ²Department of Haematology-Oncology, National University Cancer Institute, Singapore, Singapore. ³Department of Internal Medicine, University Medical Center of the Johannes Gutenberg University, Mainz, Germany. ⁴Department of Internal Medicine I, Martin-Luther-University Halle-Wittenberg, Halle/Saale, Germany. ⁵Saw Swee Hock School of Public Health, National University of Singapore, Singapore, Singapore. ⁶Institute of Medical Epidemiology, Biostatistics and Informatics, Martin-Luther-University Halle-Wittenberg, Halle/Saale, Germany

Contact: Anna Dorothea Wagner, Department of Oncology, Lausanne University Hospitals and Clinics, Rue du Bugnon 46, Lausanne, 1011, Switzerland. dorothea.wagner@chuv.ch.

Editorial group: Cochrane Upper GI and Pancreatic Diseases Group.

Publication status and date: New search for studies and content updated (conclusions changed), published in Issue 8, 2017.

Citation: Wagner AD, Syn NLX, Moehler M, Grothe W, Yong WP, Tai BC, Ho J, Unverzagt S. Chemotherapy for advanced gastric cancer. *Cochrane Database of Systematic Reviews* 2017, Issue 8. Art. No.: CD004064. DOI: [10.1002/14651858.CD004064.pub4](https://doi.org/10.1002/14651858.CD004064.pub4).

Copyright © 2017 The Cochrane Collaboration. Published by John Wiley & Sons, Ltd.

ABSTRACT

Background

Gastric cancer is the fifth most common cancer worldwide. In "Western" countries, most people are either diagnosed at an advanced stage, or develop a relapse after surgery with curative intent. In people with advanced disease, significant benefits from targeted therapies are currently limited to HER-2 positive disease treated with trastuzumab, in combination with chemotherapy, in first-line. In second-line, ramucirumab, alone or in combination with paclitaxel, demonstrated significant survival benefits. Thus, systemic chemotherapy remains the mainstay of treatment for advanced gastric cancer. Uncertainty remains regarding the choice of the regimen.

Objectives

To assess the efficacy of chemotherapy versus best supportive care (BSC), combination versus single-agent chemotherapy and different chemotherapy combinations in advanced gastric cancer.

Search methods

We searched the Cochrane Central Register of Controlled Trials, MEDLINE and Embase up to June 2016, reference lists of studies, and contacted pharmaceutical companies and experts to identify randomised controlled trials (RCTs).

Selection criteria

We considered only RCTs on systemic, intravenous or oral chemotherapy versus BSC, combination versus single-agent chemotherapy and different chemotherapy regimens in advanced gastric cancer.

Data collection and analysis

Two review authors independently identified studies and extracted data. A third investigator was consulted in case of disagreements. We contacted study authors to obtain missing information.

Main results

We included 64 RCTs, of which 60 RCTs (11,698 participants) provided data for the meta-analysis of overall survival. We found chemotherapy extends overall survival (OS) by approximately 6.7 months more than BSC (hazard ratio (HR) 0.3, 95% confidence intervals (CI) 0.24 to 0.55, 184 participants, three studies, moderate-quality evidence). Combination chemotherapy extends OS slightly (by an additional month)

versus single-agent chemotherapy (HR 0.84, 95% CI 0.79 to 0.89, 4447 participants, 23 studies, moderate-quality evidence), which is partly counterbalanced by increased toxicity. The benefit of epirubicin in three-drug combinations, in which cisplatin is replaced by oxaliplatin and 5-FU is replaced by capecitabine is unknown.

Irinotecan extends OS slightly (by an additional 1.6 months) versus non-irinotecan-containing regimens (HR 0.87, 95% CI 0.80 to 0.95, 2135 participants, 10 studies, high-quality evidence).

Docetaxel extends OS slightly (just over one month) compared to non-docetaxel-containing regimens (HR 0.86, 95% CI 0.78 to 0.95, 2001 participants, eight studies, high-quality evidence). However, due to subgroup analyses, we are uncertain whether docetaxel-containing combinations (docetaxel added to a single-agent or two-drug combination) extends OS due to moderate-quality evidence (HR 0.80, 95% CI 0.71 to 0.91, 1466 participants, four studies, moderate-quality evidence). When another chemotherapy was replaced by docetaxel, there is probably little or no difference in OS (HR 1.05; 0.87 to 1.27, 479 participants, three studies, moderate-quality evidence). We found there is probably little or no difference in OS when comparing capecitabine versus 5-FU-containing regimens (HR 0.94, 95% CI 0.79 to 1.11, 732 participants, five studies, moderate-quality evidence).

Oxaliplatin may extend (by less than one month) OS versus cisplatin-containing regimens (HR 0.81, 95% CI 0.67 to 0.98, 1105 participants, five studies, low-quality evidence). We are uncertain whether taxane-platinum combinations with (versus without) fluoropyrimidines extend OS due to very low-quality evidence (HR 0.86, 95% CI 0.71 to 1.06, 482 participants, three studies, very low-quality evidence). S-1 regimens improve OS slightly (by less than an additional month) versus 5-FU-containing regimens (HR 0.91, 95% CI 0.83 to 1.00, 1793 participants, four studies, high-quality evidence), however since S-1 is used in different doses and schedules between Asian and non-Asian population, the applicability of this finding to individual populations is uncertain.

Authors' conclusions

Chemotherapy improves survival (by an additional 6.7 months) in comparison to BSC, and combination chemotherapy improves survival (by an additional month) compared to single-agent 5-FU. Testing all patients for HER-2 status may help to identify patients with HER-2-positive tumours, for whom, in the absence of contraindications, trastuzumab in combination with capecitabine or 5-FU in combination with cisplatin has been shown to be beneficial. For HER-2 negative people, all different two- and three-drug combinations including irinotecan, docetaxel, oxaliplatin or oral 5-FU prodrugs are valid treatment options for advanced gastric cancer, and consideration of the side effects of each regimen is essential in the treatment decision. Irinotecan-containing combinations and docetaxel-containing combinations (in which docetaxel was added to a single-agent or two-drug (platinum/5-FU combination)) show significant survival benefits in the comparisons studied above. Furthermore, docetaxel-containing three-drug regimens have increased response rates, but the advantages of the docetaxel-containing three-drug combinations (DCF, FLO-T) are counterbalanced by increased toxicity. Additionally, oxaliplatin-containing regimens demonstrated a benefit in OS as compared to the same regimen containing cisplatin, and there is a modest survival improvement of S-1 compared to 5-FU-containing regimens.

Whether the survival benefit for three-drug combinations including cisplatin, 5-FU, and epirubicin as compared to the same regimen without epirubicin is still valid when second-line therapy is routinely administered and when cisplatin is replaced by oxaliplatin and 5-FU by capecitabine is questionable. Furthermore, the magnitude of the observed survival benefits for the three-drug regimens is not large enough to be clinically meaningful as defined recently by the American Society for Clinical Oncology (Ellis 2014). In contrast to the comparisons in which a survival benefit was observed by adding a third drug to a two-drug regimen at the cost of increased toxicity, the comparison of regimens in which another chemotherapy was replaced by irinotecan was associated with a survival benefit (of borderline statistical significance), but *without* increased toxicity. For this reason irinotecan/5-FU-containing combinations are an attractive option for first-line treatment. Although they need to be interpreted with caution, subgroup analyses of one study suggest that elderly people have a greater benefit from oxaliplatin, as compared to cisplatin-based regimens, and that people with locally advanced disease or younger than 65 years might benefit more from a three-drug regimen including 5-FU, docetaxel, and oxaliplatin as compared to a two-drug combination of 5-FU and oxaliplatin, a hypothesis that needs further confirmation. For people with good performance status, the benefit of second-line chemotherapy has been established in several RCTs.

PLAIN LANGUAGE SUMMARY

Chemotherapy for advanced gastric cancer

Background

Of all people with gastric cancer, in countries where screening is not routinely performed, 80% to 90% are either diagnosed at an advanced stage when the tumour is inoperable, or develop a recurrence within five years after surgery. Before starting any systemic chemotherapy in advanced disease, testing for over expression of the Human Epidermal growth factor Receptor-2 (abbreviated HER-2) testing is mandatory, and people with HER-2 over expression need, in the absence of contraindications, to be treated by a combination of a cisplatin/fluoropyrimidine-based chemotherapy and trastuzumab (i.e. a monoclonal antibody directed against the human epidermal growth factor receptor II).

Study Characteristics

Chemotherapy for advanced gastric cancer (Review)

We searched biomedical databases (MEDLINE, Embase, Cochrane Central Register of Clinical Trials) until June 2016. We included 64 RCTs, of which 60 studies with 11,698 participants contained data on overall survival, in this review. We excluded 195 studies with reasons.

Quality of the evidence

The quality of evidence ranged from very low to high, depending on the comparison and outcome being assessed. Reasons for downgrading the quality were due to risk of bias due to lack of blinded or independent radiological review, imprecision or heterogeneity.

Key results

Chemotherapy improves survival (by approximately 6.7 months) and quality of life in comparison to best supportive care alone, and first-line combination chemotherapy improves survival (by one month) compared to single-agent 5-FU.

The addition of docetaxel to platinum-fluoropyrimidine-based chemotherapy regimens appears to extend survival (by just over one additional month) at the cost of increased toxicity. Whether the benefit from adding a third drug (docetaxel or epirubicin) to a two-drug platinum-fluoropyrimidine chemotherapy combination outweighs its toxicity is unclear.

Consideration of the profile of side effects and the impact of these side effects on the individual person's quality of life, as well as the tumour burden and necessity to obtain a response rapidly is therefore essential in the choice of the regimen. Additionally, irinotecan-containing regimens prolonged overall survival (by an additional 1.6 months) compared to non-irinotecan-containing regimens.

SUMMARY OF FINDINGS

Summary of findings for the main comparison. Chemotherapy versus best supportive care for advanced gastric cancer

Chemotherapy versus best supportive care for advanced gastric cancer

Patient or population: people with advanced gastric cancer

Settings: outpatient clinics participating in international multicentre studies

Intervention: chemotherapy

Control: best supportive care alone

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	Best supportive care	Chemotherapy				
Overall survival	Study population		HR 0.37 (0.24 to 0.55)	184 (3 studies)	⊕⊕⊕⊖ moderate ¹	Weighted average of median survival durations from included studies
	4.3 months	11.0 months				
Time to progression	Study population		HR 0.31 (0.22 to 0.43)	144 (2 studies)	⊕⊕⊕⊖ moderate ¹	Weighted average of median survival durations from included studies
	2.5 months	7.4 months				

*For time-to-event outcomes, e.g. overall survival, the **assumed** and **corresponding risks** were obtained by calculating the weighted average of the median survival durations reported in included studies. For dichotomous outcomes, the **assumed** and **corresponding risks** (and their 95% confidence interval) are based on proportions of events in the control and intervention groups respectively.

CI: Confidence interval; **HR:** Hazard ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Early termination of [Pyrhönen 1995](#); downgraded by one level for risk of bias.

Outcomes shown include those which were measured in the studies, or reported in a consistent fashion across included studies. Several critical outcomes (e.g. tumour response, treatment-related death, and discontinuation due to toxicity) were not evaluated or reported in a consistent fashion in these studies, as they were mainly conducted before year 2000.

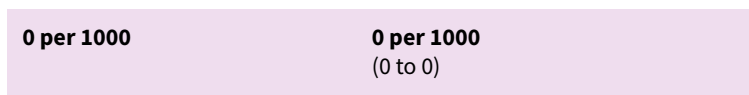
Summary of findings 2. Combination versus single-agent chemotherapy for advanced gastric cancer
Combination versus single-agent chemotherapy for advanced gastric cancer
Patient or population: people with advanced gastric cancer

Settings: outpatient clinics participating in international multicentre studies

Intervention: combination

Control: single-agent chemotherapy

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	Single-agent chemotherapy	Combination				
Overall survival	Study population		HR 0.84 (0.79 to 0.89)	4447 (23)	⊕⊕⊕○ moderate ¹	Weighted average of median survival durations from included studies
	<ul style="list-style-type: none"> 10.5 months in studies published after year 2000 6.4 months in studies published before year 2000 	<ul style="list-style-type: none"> 11.6 months in studies published after year 2000 7.3 months in studies published before year 2000 				
Tumour response	Study population		OR 2.30 (1.94 to 2.72)	2833 (18)	⊕⊕⊕⊕ high ¹	
	226 per 1000	402 per 1000 (361 to 442)				
	Moderate					
	231 per 1000	409 per 1000 (368 to 450)				
Time to progression	Study population		HR 0.69 (0.55 to 0.87)	720 (4)	⊕⊕⊕○ moderate ¹	Weighted average of median survival durations from included studies
	2.8 months	4.1 months				
Treatment-related death	Study population		OR 1.64 (0.83 to 3.24)	3876 (18)	⊕⊕○○ moderate ²	
	5 per 1000	9 per 1000 (4 to 17)				
	Moderate					



*For time-to-event outcomes, e.g. overall survival, the **assumed** and **corresponding risks** were obtained by calculating the weighted average of the median survival durations reported in included studies. For dichotomous outcomes, the **assumed** and **corresponding risks** (and their 95% confidence interval) are based on proportions of events in the control and intervention groups respectively.

CI: Confidence interval; **OR:** Odds ratio; **HR:** Hazard ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Downgraded by one level for risk of bias.

² Downgraded by two levels for serious imprecision.

Summary of findings 3. 5-FU/cisplatin/anthracycline combinations versus 5-FU/cisplatin combinations (without anthracyclines) for advanced gastric cancer

5-FU/cisplatin/anthracycline combinations versus 5-FU/cisplatin combinations (without anthracyclines) for advanced gastric cancer

Patient or population: people with advanced gastric cancer

Settings: outpatient clinics participating in international multicentre studies

Intervention: 5-FU/cisplatin/anthracycline combinations

Control: 5-FU/cisplatin combinations (without anthracyclines)

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	5-FU/cisplatin combinations (without anthracyclines)	5-FU/cisplatin/anthracycline combinations				
Overall survival	Study population		HR 0.74 (0.61 to 0.89)	579 (4)	⊕⊕⊕⊖ moderate ¹	Weighted average of median survival durations from included studies
	8.6 months	9.9 months				
Tumour response	Study population		OR 2.86 (1.14 to 7.16)	78 (1)	⊕⊕⊖⊖ low ²	

	385 per 1000	641 per 1000 (416 to 817)			
	Moderate				
	385 per 1000	642 per 1000 (416 to 818)			
Time to progression	Study population		HR 0.62 (0.38 to 0.98)	78 (1)	⊕⊕⊕⊕ low ²
	7.9 months	12.1 months			Median survival durations from the only included study

*For time-to-event outcomes, e.g. overall survival, the **assumed** and **corresponding risks** were obtained by calculating the weighted average of the median survival durations reported in included studies. For dichotomous outcomes, the **assumed** and **corresponding risks** (and their 95% confidence interval) are based on proportions of events in the control and intervention groups respectively.

CI: Confidence interval; **OR:** Odds ratio; **HR:** Hazard ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Downgraded by one level for risk of bias.

² Downgraded by two levels for serious imprecision.

Outcomes shown include those which were measured in the studies, or reported in a consistent fashion across included studies. Several critical outcomes (e.g. treatment-related death and discontinuation due to toxicity) were not evaluated or reported in a consistent fashion in these studies, as they were mainly conducted before year 2000.

Summary of findings 4. 5-FU/cisplatin/anthracycline combinations versus 5-FU/anthracycline combinations (without cisplatin) for advanced gastric cancer

5-FU/cisplatin/anthracycline combinations versus 5-FU/anthracycline combinations (without cisplatin) for advanced gastric cancer

Patient or population: people with advanced gastric cancer

Settings: outpatient clinics participating in international multicentre studies

Intervention: 5-FU/cisplatin/anthracycline combinations

Control: 5-FU/cisplatin combinations (without anthracyclines)

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				

	5-FU/anthracycline combinations (without cisplatin)	5-FU/cisplatin/anthracycline combinations				
Overall survival	Study population		HR 0.82 (0.73 to 0.92)	1147 (7)	⊕⊕⊕⊖ low 1,2	Weighted average of median survival durations from included studies
	6.2 months	8.4 months				

*For time-to-event outcomes, e.g. overall survival, the **assumed** and **corresponding risks** were obtained by calculating the weighted average of the median survival durations reported in included studies. For dichotomous outcomes, the **assumed** and **corresponding risks** (and their 95% confidence interval) are based on proportions of events in the control and intervention groups respectively.

CI: Confidence interval; **HR:** Hazard ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Downgraded by one level for risk of bias.

² Downgraded by one level for statistical heterogeneity.

Several critical outcomes (i.e. tumour response, progression-free survival, treatment-related death and discontinuation due to toxicity) were not evaluated or reported in a consistent fashion in these studies, most of which were conducted before year 2000.

Summary of findings 5. Irinotecan versus non-irinotecan-containing regimens for advanced gastric cancer

Irinotecan versus non-irinotecan-containing regimens for advanced gastric cancer

Patient or population: people with advanced gastric cancer

Settings: outpatient clinics participating in international multicentre studies

Intervention: irinotecan

Control: non-irinotecan-containing regimens

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	Non-irinotecan-containing regimens	Chemotherapy with Irinotecan				
Overall survival	Study population		HR 0.87 (0.80 to 0.95)	2135 (10 studies)	⊕⊕⊕⊖ moderate ¹	Weighted average of median survival durations from included studies

	9.7 months	11.3 months				
Overall survival - Substitutive comparisons	Study population		HR 0.87	826 (6 studies)	⊕⊕⊕⊖ moderate ¹	Weighted average of median survival durations from included studies
	9.1 months	9.9 months	(0.75 to 1.00)			
Overall survival - Additive comparisons	Study population		HR 0.88	500 (3 studies)	⊕⊕⊕⊖ low ^{1,2}	Weighted average of median survival durations from included studies
	10.9 months	11.9 months	(0.76 to 1.03)			
Overall survival - Other comparisons	Study population		HR 0.87	809 (2 studies)	⊕⊕⊕⊖ very low ^{1,3}	Weighted average of median survival durations from included studies
	11.4 months	12.6 months	(0.76 to 1.00)			
Tumour response	Study population		OR 1.72	1266 (10 studies)	⊕⊕⊕⊖ low ³	
	288 per 1000	410 per 1000 (334 to 493)	(1.24 to 2.40)			
	Moderate					
	275 per 1000	395 per 1000 (320 to 477)				
Tumour response - Substitutive comparisons	Study population		OR 1.53	756 (6 studies)	⊕⊕⊕⊖ low ³	
	297 per 1000	393 per 1000 (282 to 514)	(0.93 to 2.50)			
	Moderate					
	294 per 1000	389 per 1000 (279 to 510)				
Tumour response - Additive comparisons	Study population		OR 2.18	345 (3 studies)	⊕⊕⊕⊖ low ^{1,2}	
	224 per 1000	386 per 1000 (265 to 522)	(1.25 to 3.80)			
	Moderate					
	219 per 1000	379 per 1000 (260 to 516)				

Tumour response - Other comparisons	Study population	OR 1.87 (0.89 to 3.91)	165 (2 studies)	⊕⊕⊕⊕ very low 1,2,4	
	376 per 1000 530 per 1000 (350 to 702)				
	Moderate				
	367 per 1000 520 per 1000 (340 to 694)				
Progression-free survival	Study population	HR 0.76 (0.69 to 0.84)	1640 (7 studies)	⊕⊕⊕⊕ high	Weighted average of median survival durations from included studies
	4.4 months 5.9 months				
Progression-free survival - Substitutive comparison	Study population	HR 0.85 (0.72 to 1.00)	741 (5 studies)	⊕⊕⊕⊖ moderate 1	Weighted average of median survival durations from included studies
	4.2 months 5.3 months				
Progression-free survival - Additive comparisons	Study population	HR 0.51 (0.33 to 0.77)	90 (1)	⊕⊕⊕⊖ moderate 2	Median survival durations from the only included study
	3.2 months 6.9 months				
Progression-free survival - Other comparisons	Study population	HR 0.74 (0.66 to 0.84)	809 (2 studies)	⊕⊕⊕⊕ high	Weighted average of median survival durations from included studies
	5.4 months 6.6 months				
Treatment-related death	Study population	OR 0.88 (0.23 to 3.32)	1979 (9 studies)	⊕⊕⊕⊖ low 2,4	
	10 per 1000 9 per 1000 (2 to 32)				
	Moderate				
	2 per 1000 2 per 1000 (0 to 7)				
Treatment discontinuation due to toxicity	Study population	OR 1.00 (0.46 to 2.20)	1979 (9 studies)	⊕⊕⊕⊖ very low 2,3	
	137 per 1000 137 per 1000 (68 to 258)				
	Moderate				
	215 per 1000 215 per 1000				

(112 to 376)

*For time-to-event outcomes, e.g. overall survival, the **assumed** and **corresponding risks** were obtained by calculating the weighted average of the median survival durations reported in included studies. For dichotomous outcomes, the **assumed** and **corresponding risks** (and their 95% confidence interval) are based on proportions of events in the control and intervention groups respectively.

CI: Confidence interval; **OR:** Odds ratio; **HR:** Hazard ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Downgraded by one level for risk of bias.

² Downgraded by one level for imprecision.

³ Downgraded by two levels for severe statistical heterogeneity.

⁴ Downgraded by one level for statistical heterogeneity.

Summary of findings 6. Docetaxel versus non-docetaxel-containing regimens for advanced gastric cancer

Docetaxel versus non-docetaxel-containing regimens for advanced gastric cancer

Patient or population: people with advanced gastric cancer

Settings: outpatient clinics participating in international multicentre studies

Intervention: docetaxel

Control: non-docetaxel-containing regimens

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	Non-docetaxel-containing regimens	Chemotherapy with docetaxel				
Overall survival	Study population		HR 0.86 (0.78 to 0.95)	2001 (8 studies)	⊕⊕⊕⊕ high	Weighted average of median survival durations from included studies
	9.9 months	11.2 months				
Overall survival - Substitutive comparisons	Study population		HR 1.05 (0.87 to 1.27)	479 (3 studies)	⊕⊕⊕⊖ moderate ¹	Weighted average of median survival durations from included studies
	9.4 months	9.2 months				

Overall survival - Additive comparisons	Study population	HR 0.80	1466 (4 studies)	⊕⊕⊕⊖ moderate ²	Weighted average of median survival durations from included studies
	10.6 months 12.3 months	(0.71 to 0.91)			
Overall survival - Other comparisons	Study population	HR 0.80	56 (1 study)	⊕⊖⊖⊖ very low ^{1,2,3}	Median survival durations from the only included study
	9.5 months 11.9 months	(0.46 to 1.39)			
Tumour response	Study population	OR 1.37	1820 (9 studies)	⊕⊕⊕⊖ moderate ⁴	
	311 per 1000 382 per 1000 (317 to 452)	(1.03 to 1.83)			
	Moderate				
	310 per 1000 381 per 1000 (316 to 451)				
Tumour response - Substitutive comparison	Study population	OR 1.03	525 (4 studies)	⊕⊕⊕⊖ moderate ¹	
	314 per 1000 320 per 1000 (245 to 407)	(0.71 to 1.50)			
	Moderate				
	327 per 1000 334 per 1000 (256 to 422)				
Tumour response - Additive comparison	Study population	OR 1.83	1235 (4 studies)	⊕⊕⊕⊕ high	
	295 per 1000 434 per 1000 (378 to 493)	(1.45 to 2.32)			
	Moderate				
	296 per 1000 435 per 1000 (379 to 494)				
Tumour response - Other comparison	Study population	OR 0.33	60 (1 study)	⊕⊖⊖⊖ very low ^{1,3}	
	600 per 1000 331 per 1000 (153 to 590)	(0.12 to 0.96)			

	Moderate					
	600 per 1000	331 per 1000 (153 to 590)				
Time to progression	Study population		HR 1.06	360 (2 studies)	⊕⊕⊕⊕ very low 1,2,3	Weighted average of median survival durations from included studies
	6.0 months	5.9 months	(0.85 to 1.32)			
Progression-free survival	Study population		HR 0.76	1498 (5 studies)	⊕⊕⊕⊕ moderate 4	Weighted average of median survival durations from included studies
	4.8 months	6.0 months	(0.63 to 0.91)			
Progression-free survival - Substitutive comparisons	Study population		HR 1.15	119 (1 study)	⊕⊕⊕⊕ very low 1,2,3	Median survival durations from the only included study
	4.9 months	4.6 months	(0.77 to 1.72)			
Progression-free survival - Additive comparison	Study population		HR 0.70	1323 (3 studies)	⊕⊕⊕⊕ high	Weighted average of median survival durations from included studies
	4.3 months	6.0 months	(0.61 to 0.81)			
Progression-free survival - Other comparison	Study population		HR 0.94	56 (1 study)	⊕⊕⊕⊕ very low 1,3	Median survival durations from the only included study
	6.4 months	6.8 months	(0.55 to 1.60)			
Treatment-related death	Study population		OR 1.10	2113 (7 studies)	⊕⊕⊕⊕ moderate 1	
	12 per 1000	14 per 1000 (7 to 27)				
	Moderate					
	5 per 1000	5 per 1000 (3 to 11)				
Treatment discontinuation due to toxicity	Study population		OR 0.81	1066 (5 studies)	⊕⊕⊕⊕ low 1,4	
	211 per 1000	178 per 1000 (124 to 251)		(0.53 to 1.25)		
	Moderate					
	197 per 1000	166 per 1000				

(115 to 235)

*For time-to-event outcomes, e.g. overall survival, the **assumed** and **corresponding risks** were obtained by calculating the weighted average of the median survival durations reported in included studies. For dichotomous outcomes, the **assumed** and **corresponding risks** (and their 95% confidence interval) are based on proportions of events in the control and intervention groups respectively.

CI: Confidence interval; **OR:** Odds ratio; **HR:** Hazard ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Downgraded by one level for imprecision.

² Downgraded by one level for risk of bias.

³ Downgraded by two levels for serious imprecision.

⁴ Downgraded by one level for statistical heterogeneity.

Summary of findings 7. Capecitabine versus 5-FU-containing regimens for advanced gastric cancer

Capecitabine versus 5-FU-containing regimens for advanced gastric cancer

Patient or population: people with advanced gastric cancer

Settings: outpatient clinics participating in international multicentre studies with approximately half of all participants enrolled from Asian countries

Intervention: capecitabine

Control: 5-FU-containing regimens

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	5-FU-containing regimens	Capecitabine-containing regimens				
Overall Survival	Study population		HR 0.94 (0.79 to 1.11)	732 (5 studies)	⊕⊕⊕⊖ moderate ¹	Weighted average of median survival durations from included studies
	10.9 months	10.8 months				
Tumour response	Study population		OR 0.85 (0.40 to 1.79)	636 (4 studies)	⊕⊖⊖⊖ very low ^{2,3}	
	384 per 1000	347 per 1000				

		(200 to 528)			
	Moderate				
	394 per 1000	356 per 1000 (206 to 538)			
Time to progression	Study population		HR 0.72	85 (1 study)	⊕⊕⊕⊕ very low ^{1,3}
	5.5 months	6.8 months	(0.47 to 1.12)		Median survival durations from the only included study
Progression-free survival	Study population		HR 0.98	647 (4 studies)	⊕⊕⊕⊕ very low ^{1,3,4}
	6.7 months	6.5 months	(0.77 to 1.23)		Weighted average of median survival durations from included studies
Treatment-related death	Study population		OR 1.88	481 (2 studies)	⊕⊕⊕⊕ very low ^{1,2,3}
	21 per 1000	38 per 1000 (5 to 241)	(0.23 to 15.15)		
	Moderate				
	24 per 1000	44 per 1000 (6 to 271)			
Treatment discontinuation due to toxicity	Study population		OR 0.99	311 (1 study)	⊕⊕⊕⊕ low ³
	181 per 1000	179 per 1000 (110 to 281)	(0.56 to 1.77)		
	Moderate				
	181 per 1000	180 per 1000 (110 to 281)			

*For time-to-event outcomes, e.g. overall survival, the **assumed** and **corresponding risks** were obtained by calculating the weighted average of the median survival durations reported in included studies. For dichotomous outcomes, the **assumed** and **corresponding risks** (and their 95% confidence interval) are based on proportions of events in the control and intervention groups respectively.

CI: Confidence interval; **OR:** Odds ratio; **HR:** Hazard ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

- ¹ Downgraded by one level for risk of bias.
- ² Downgraded by two levels for severe statistical heterogeneity.
- ³ Downgraded by two levels for serious imprecision.
- ⁴ Downgraded by one level for statistical heterogeneity.

Summary of findings 8. Oxaliplatin versus the same regimen including cisplatin for advanced gastric cancer

Oxaliplatin versus the same regimen including cisplatin for advanced gastric cancer

Patient or population: people with advanced gastric cancer

Settings: outpatient clinics participating in international multicentre studies with the majority of participants enrolled in Asia

Intervention: oxaliplatin-containing regimen

Control: the same regimen including cisplatin

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	Cisplatin-containing regimen	Oxaliplatin-containing regimen				
Overall Survival	Study population		HR 0.81 (0.67 to 0.98)	1105 (5 studies)	⊕⊕⊕⊖ low ^{1,2}	Weighted average of median survival durations from included studies
	11.3 months	14.0 months				
Tumour response	Study population		OR 1.38 (1.08 to 1.76)	1081 (5 studies)	⊕⊕⊕⊕ moderate ¹	
	468 per 1000	548 per 1000 (487 to 607)				
	Moderate					
	458 per 1000	538 per 1000 (477 to 598)				
Progression-free survival	Study population		HR 0.88 (0.66 to 1.19)	1034 (4 studies)	⊕⊕⊕⊖ low ^{1,3}	Weighted average of median survival dura-
	4.9 months	6.0 months				

tions from included studies

Treatment-related death	Study population	OR 0.47	1132 (5 studies)	⊕⊕⊕⊖ low 1,3
	20 per 1000	9 per 1000 (3 to 25)	(0.17 to 1.30)	
	Moderate			
Treatment discontinuation due to toxicity	Study population	OR 0.97	970 (3 studies)	⊕⊕⊕⊖ very low 1,2,3
	95 per 1000	93 per 1000 (44 to 183)	(0.44 to 2.13)	
	Moderate			
	102 per 1000	99 per 1000 (48 to 195)		

*For time-to-event outcomes, e.g. overall survival, the **assumed** and **corresponding risks** were obtained by calculating the weighted average of the median survival durations reported in included studies. For dichotomous outcomes, the **assumed** and **corresponding risks** (and their 95% confidence interval) are based on proportions of events in the control and intervention groups respectively.

CI: Confidence interval; **OR:** Odds ratio; **HR:** Hazard ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Downgraded by one level for risk of bias.

² Downgraded by one level for statistical heterogeneity.

³ Downgraded by one level for imprecision.

Summary of findings 9. Taxane-platinum-fluoropyrimidine combinations versus taxane-platinum (without fluoropyrimidine) for advanced gastric cancer

Taxane-platinum-fluoropyrimidine combinations versus taxane-platinum (without fluoropyrimidine) for advanced gastric cancer

Patient or population: people with advanced gastric cancer
Settings: outpatient clinics participating in international multicentre studies, without Asian representation
Intervention: taxane-platinum-fluoropyrimidine combinations
Control: taxane-platinum (without fluoropyrimidine)

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	Taxane-platinum (without fluoropyrimidine)	Taxane-platinum-fluoropyrimidine combination				
Overall survival	Study population		OR 0.86 (0.71 to 1.06)	482 (3 studies)	⊕⊕⊕⊕ very low ^{1,2}	Weighted average of median survival durations from included studies
	10.0 months	11.7 months				
Tumour response	Study population		OR 2.08 (1.37 to 3.15)	482 (3 studies)	⊕⊕⊕⊕ low ^{1,3}	
	234 per 1000	389 per 1000 (295 to 491)				
	Moderate					
	231 per 1000	385 per 1000 (292 to 486)				
Progression-free survival	Study population		OR 0.74 (0.59 to 0.93)	482 (3 studies)	⊕⊕⊕⊕ moderate ¹	Weighted average of median survival durations from included studies
	4.4 months	5.7 months				
Treatment-related death	Study population		OR 1.95 (0.73 to 5.17)	482 (3 studies)	⊕⊕⊕⊕ very low ^{1,4}	
	26 per 1000	50 per 1000 (19 to 121)				
	Moderate					
	13 per 1000	25 per 1000 (10 to 64)				

Treatment discontinuation due to toxicity	Study population		OR 1.71 (0.79 to 3.69)	234 (2 studies)	⊕○○○ very low ^{1,4}
	105 per 1000	167 per 1000 (85 to 303)			
	Moderate				
	99 per 1000	158 per 1000 (80 to 288)			

*For time-to-event outcomes, e.g. overall survival, the **assumed** and **corresponding risks** were obtained by calculating the weighted average of the median survival durations reported in included studies. For dichotomous outcomes, the **assumed** and **corresponding risks** (and their 95% confidence interval) are based on proportions of events in the control and intervention groups respectively.

CI: Confidence interval; **OR:** Odds ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Downgraded by one level for risk of bias.

² Downgraded by two levels for severe statistical heterogeneity.

³ Downgraded by one level for imprecision.

⁴ Downgraded by two levels for serious imprecision.

Summary of findings 10. S-1 versus 5-FU-containing regimens for advanced gastric cancer

S-1 versus 5-FU-containing regimens for advanced gastric cancer

Patient or population: people with advanced gastric cancer

Settings: outpatient clinics participating in international multicentre studies, mostly performed in Asia

Intervention: S-1-containing regimens

Control: 5-FU-containing regimens

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	5-FU-containing regimens	S-1 containing regimens				

Overall Survival	Study population	HR 0.91 (0.83 to 1.00)	1793 (4 studies)	⊕⊕⊕⊕ high	Weighted average of median survival durations from included studies
	9.1 months 9.6 months				
Tumour response	Study population	OR 1.73 (1.01 to 2.94)	1753 (7 studies)	⊕⊕⊕⊕ very low ^{1,2}	
	256 per 1000 374 per 1000 (258 to 503)				
	Moderate				
	320 per 1000 449 per 1000 (322 to 580)				
Progression-free survival	Study population	HR 0.85 (0.70 to 1.04)	1942 (4 studies)	⊕⊕⊕⊕ low ¹	Weighted average of median survival durations from included studies
	4.3 months 5.0 months				
Time-to treatment failure	Study population	HR 0.88 (0.76 to 1.01)	1818 (5 studies)	⊕⊕⊕⊕ low ¹	Weighted average of median survival durations from included studies
	3.1 months 3.9 months				
Treatment-related deaths	Study population	OR 0.56 (0.30 to 1.06)	1962 (4 studies)	⊕⊕⊕⊕ moderate ²	
	27 per 1000 15 per 1000 (8 to 28)				
	Moderate				
	5 per 1000 3 per 1000 (2 to 5)				
Treatment discontinuation due to toxicity	Study population	OR 0.85 (0.63 to 1.13)	1726 (3 studies)	⊕⊕⊕⊕ high	
	128 per 1000 111 per 1000 (85 to 142)				
	Moderate				
	144 per 1000 125 per 1000 (96 to 160)				

*For time-to-event outcomes, e.g. overall survival, the **assumed** and **corresponding risks** were obtained by calculating the weighted average of the median survival durations reported in included studies. For dichotomous outcomes, the **assumed** and **corresponding risks** (and their 95% confidence interval) are based on proportions of events in the control and intervention groups respectively.

CI: Confidence interval; **OR:** Odds ratio; **HR:** Hazard ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Downgraded by two levels for severe statistical heterogeneity.

² Downgraded by one level for imprecision.

BACKGROUND

Description of the condition

Epidemiology and pathogenesis

With an estimated 1.3 million new cases in 2015, gastric cancer is currently the fifth most common malignancy and the third leading cause of cancer-related mortality worldwide (GBD Cancer Collaboration 2017). Only approximately 25% of all people with gastric cancer have resectable disease at presentation. Stomach cancer incidence rates show substantial variation internationally, with endemic regions in Asia, Eastern Europe and South America (Ferro 2014). *Helicobacter pylori*, atrophic gastritis, intestinal metaplasia, and dysplasia have been identified as important steps in the pathogenesis of gastric cancer (Correa 1996). Due to improvements in food conservation and diet, as well as eradication of *Helicobacter pylori*, gastric cancer incidence and mortality has steadily fallen in the last 50 years (Peleteiro 2012).

In contrast, a dramatic rise in cardiac and gastroesophageal junction tumour incidence rates has been observed in middle-aged, male, white Caucasians (Abrams 2013; Sharma 2003; Wu 2001). A proportion of these cases seems to be associated with Barrett's epithelium (intestinal metaplasia of the distal oesophagus), developing from chronic oesophageal reflux disease (MacDonald 1992; Wu-Williams 1990). Although it is difficult to determine whether these cancers are gastroesophageal junction tumours or distal oesophageal malignancies (Rusch 2004), in clinical studies for advanced disease they are usually treated in the same manner.

Gastric cancer is a heterogenous disease entity, with major differences in growth patterns, differentiation, and molecular pathogenesis. More than 90% of stomach tumours are adenocarcinomas. While Lauren already in 1965 distinguished (Lauren 1965) the *well-differentiated or intestinal type* and the *undifferentiated or diffuse-type*, the current World Health Organization (WHO) classification Bosman 2010 differentiates the following five major histopathological subtypes: papillary, tubular, and mucinous adenocarcinoma, as well as poorly cohesive (with or without signet cells) and mixed carcinoma. While the first three types correspond to the former "well differentiated or intestinal type", the undifferentiated or diffuse type according to Lauren corresponds to the poorly cohesive type in the current classification.

While the intestinal type is more common in males, older age groups, and in high-risk geographic areas, diffuse-type carcinomas have a more equal male to female distribution, are more frequent in younger individuals, and have a more uniform geographic distribution (Crew 2004; Kelley 2003; Lauren 1965; Munoz 1968). Ninety per cent of gastric cancers are sporadic. Hereditary diffuse gastric cancer is rare, with less than 3% of cases. According to a recently published landmark paper (TCGA 2014), which describes the results of a comprehensive molecular evaluation of 295 primary gastric adenocarcinomas as part of The Cancer Genome Atlas (TCGA), the following four molecular subtypes can be distinguished.

1. Tumours positive for Epstein–Barr virus, which display recurrent PIK3CA mutations, extreme DNA hyper methylation, and amplification of JAK2, CD274 (also known as PD-L1) and PDCD1LG2 (also known as PD-L2).

2. Microsatellite unstable tumours, which show elevated mutation rates.
3. Genomically stable tumours, which are enriched for the diffuse histological variant and mutations of RHOA or fusions involving RHO-family GTPase-activating proteins.
4. Tumours with chromosomal instability, which show marked aneuploidy and focal amplification of receptor tyrosine kinases.

While this classification has no impact in the choice of systemic treatment at present, it will provide a roadmap for patient stratification and development of targeted therapies in the future.

In contrast, the over expression of the Human Epidermal growth factor Receptor-2 (HER-2), which is observed in 10% to 20% of the people, is clinically relevant today as it predicts a significant benefit from treatment with trastuzumab (Bang 2010). Further details of the pathogenesis of gastric cancer have been reviewed recently by Wadhwa and colleagues (Wadhwa 2013).

Prognosis and management options

Apart from endoscopic treatment for a minority of very small tumours, partial or complete gastrectomy with lymphadenectomy is the only potentially curative therapy for gastric cancer. Stage I to IV M0 tumours are principally resectable (MacDonald 2001a). However, although surgery carries a high cure rate for stage IA and IB cancers, the results for stage IIIA and IIIB cancers are poor. Many people with advanced disease, especially stage IIIA/B, are technically inoperable. Results for both resectable and locally advanced gastric cancer may be improved by either perioperative (e.g. Cunningham 2006; Ychou 2011) or adjuvant chemotherapy (Bang 2012; Paoletti 2013; Sakuramoto 2007). Unfortunately, even after an apparently 'curative' gastrectomy, relapse rates in prospective studies remain in the range of 40% to 60% (Bonenkamp 1999; Cunningham 2006; MacDonald 2001b; Songun 2010) in European studies. In the Western world, most people are diagnosed at an advanced stage, when the tumour is inoperable. People with inoperable, recurrent or metastatic tumours have a poor prognosis with a median survival time of three to five months without chemotherapy. Several small randomised studies have provided evidence that first-line chemotherapy improves survival in these people (Glimelius 1994; Pyrhönen 1995; Scheithauer 1996), but benefit has to be weighed against treatment-related toxicities. Furthermore, second-line chemotherapy has shown to improve survival and quality of life in several recent randomised studies (Ford 2014; Kang 2012; Thuss-Patience 2011).

While a significant number of phase-III studies have studied the value of targeted therapies in advanced gastric cancer (e.g. Lordick 2013; Ohtsu 2011; Ohtsu 2013; Satoh 2014), only three phase III studies (Bang 2010; Fuchs 2014; Wilke 2014) have had positive results and impact on clinical practice:

According to the randomised phase III "TOGA" study (Bang 2010), response rate, progression-free- and overall survival are greatly improved by adding the monoclonal antibody trastuzumab to the combination of cisplatin and capecitabine in HER-2 positive gastric cancer, and introduced trastuzumab as a standard of care for HER-2 positive disease. This study changed the workup of all people with advanced gastric cancer since all people with advanced gastric cancer must now undergo HER-2 testing before the initiation of any chemotherapy and, in the absence of contraindications, be treated with trastuzumab in combination with cisplatin and 5-FU

or capecitabine in case of HER-2 over expression on IHC (IHC 3+, or IHC 2+/FISH+). All other treatment options discussed in this review (oxaliplatin, irinotecan, docetaxel, as well as the anthracycline-containing regimen ECF) are therefore valid only for people with HER-2 negative disease. Studies on combinations of other chemotherapies with trastuzumab are currently limited to phase II and cannot be recommended for this reason. Furthermore, the VEGF-receptor-targeting antibody ramucirumab, with or without chemotherapy, has been evaluated in two phase III studies as second-line treatment, and emerged as a new treatment option in this indication. Thus, despite this recent progress, chemotherapy remains the mainstay of treatment for the majority of people with advanced gastric cancer.

Description of the intervention

Systemic chemotherapy

5-FU is not only the most important and extensively studied single agent in this disease, but it is part of most combination chemotherapy regimens as well. Its single-agent response rate is about 20%. Differences in effect and toxicity profile are the reasons for its application as continuous infusion. Oral capecitabine (Cunningham 2008) or S-1 (Ajani 2010) may replace infusional 5-FU, thus avoiding the risk and inconvenience associated with portable pumps. Other single agents with relevant activities are cisplatin (Leichman 1991) and anthracyclines (Preusser 1988). Furthermore, oxaliplatin, docetaxel, and irinotecan have been evaluated in recent phase III studies (Al Batran 2008; Cunningham 2008; Dank 2008; Van Cutsem 2006).

How the intervention might work

5-FU, an antimetabolite, pyrimidine-antagonist and inhibitor of thymidilate-synthetase is the backbone of chemotherapy in gastric cancer. Capecitabine is an oral fluoropyrimidine that is selectively activated in tumour tissue by a three-step enzymatic conversion, S-1 another oral fluoropyrimidine. Cisplatin is an alkylating agent, which acts through induction of intra- and inter-strand crosslinks. The diamino-cyclohexane platinum derivative oxaliplatin also leads to the formation of DNA crosslinks, but they are not recognised by the intracellular mismatch repair system. Docetaxel, as well as paclitaxel are agents which inhibit depolymerization of micro tubuli. Irinotecan is an inhibitor of topoisomerase I, thereby disrupting DNA replication and cell division. These drugs have major differences in their toxicity profile.

Why it is important to do this review

Combination chemotherapy has become an accepted standard for first-line treatment. Although a large number of different regimens have been tested in randomised studies, uncertainty remains regarding the choice of the regimen.

OBJECTIVES

To evaluate the effect of chemotherapy in participants with advanced adenocarcinoma of the stomach and gastroesophageal junction.

Comparisons were as follows.

1. First-line chemotherapy plus best supportive care (BSC) versus BSC alone.

2. First-line combination versus single-agent chemotherapy.
3. First-line 5-FU/cisplatin/anthracycline-containing combinations versus 5-FU/cisplatin combinations (without anthracyclines).
4. First-line 5-FU/cisplatin/anthracycline-containing combinations versus 5-FU/anthracycline combinations (without cisplatin).
5. First-line chemotherapy with irinotecan versus non-irinotecan-containing regimens.
6. First-line chemotherapy with docetaxel versus non-docetaxel-containing regimens.
7. First-line chemotherapy with capecitabine versus 5-FU-containing regimens.
8. First-line chemotherapy with oxaliplatin versus the same regimen containing cisplatin.
9. First-line taxane-platinum-fluoropyrimidine combinations versus taxane-platinum (without fluoropyrimidine).
10. First-line S-1 versus 5-FU-containing regimens.

Due to limited information, we considered second-line therapy only in selected sensitivity analyses of first-line therapy where data were available. In addition to comparisons 1 and 2, which were planned and described in the first version of the protocol, we performed two more comparisons (3 and 4) in the original version of the review (Wagner 2005). As there was a large number of categories of different combination chemotherapy regimens and the number of relevant studies in each category was not known when writing the protocol, it was impossible to plan in advance the best way to compare directly the different categories of combination chemotherapies. We chose to perform these additional comparisons based on their clinical relevance and the availability of a sufficient number of relevant studies.

In the first update of this review (Other published versions of this review, Wagner 2010), comparisons (5) to (8) (5: First-line chemotherapy with irinotecan versus non-irinotecan-containing regimens, 6: First-line chemotherapy with docetaxel versus non-docetaxel-containing regimens, 7: First-line chemotherapy with capecitabine versus 5-FU-containing regimens, 8: First-line chemotherapy with oxaliplatin versus the same regimen containing cisplatin) were added. In this second update, comparisons (9) to (10) (9: First-line taxane-platinum-fluoropyrimidine combinations versus taxane-platinum (without fluoropyrimidine) and 10: First-line S-1 versus 5-FU-containing regimens) were added (see - Differences between protocol and review).

METHODS

Criteria for considering studies for this review

Types of studies

We included only randomised controlled studies, with or without blinding. We included abstracts or unpublished data if sufficient information on study design, characteristics of participants, interventions, and outcomes was available and if full information and final results were confirmed by the first author. We excluded cross-over studies in order to assess the overall treatment effect on survival. We excluded quasi-randomised studies, e.g. treatment allocation alternate or by date of birth, as we considered this study design to be not of sufficiently high quality.

Types of participants

We included participants with histologically confirmed, unresectable (as decided by a multidisciplinary team), recurrent or metastatic adenocarcinoma of the stomach or gastroesophageal junction without any prior chemotherapy or radiotherapy for comparisons 1 to 10. We also included studies which included participants with adenocarcinoma of the distal oesophagus. Most studies included participants with locally advanced, relapsed and/or metastatic tumours, with the greater number of participants already having metastatic disease. However, in some studies only participants with locally advanced cancer of the stomach were reported in order to assess secondary resectability. We did not consider these studies in this review. The proportion of participants with locally advanced versus metastatic tumours is given for each study.

Types of interventions

We included studies of systemic intravenous or oral, first-line chemotherapy and/or best supportive care (BSC). Chemotherapy encompasses all cytotoxic or anti-neoplastic drug treatment, but excluding hormonal, biological, or targeted therapies, which are the subject of a separate Cochrane review (Song 2016). However, studies on targeted therapies with clinical impact, as well as studies on second-line chemotherapy are considered in the discussion.

We included single-agent as well as combination chemotherapy studies in all doses and schedules, but did not consider combined radio-chemotherapy.

Types of outcome measures

Primary outcomes

1. Overall survival on intention-to-treat analysis. Median, one-, two- and three-year as well as five-year survival in participants with locally advanced, secondary resectable tumours.

Secondary outcomes

1. Tumour response.
2. Time to progression.
3. Secondary resectability in participants with locally advanced gastric cancer.
4. Toxicity, classified according to WHO or National Cancer Institute Common Toxicity Criteria (NCI-CTC).

Quality of life is difficult to measure and was assessed with various instruments. Quality of life results of recent phase-III studies are described in the results section and considered in the discussion if available.

Search methods for identification of studies

Electronic searches

We originally identified studies by searching the Cochrane Central Register of Controlled Trials (CENTRAL) (the Cochrane Library 2004, Issue 1), MEDLINE, and Embase up to February 2004 and reference lists of articles. We also contacted pharmaceutical companies as well as national and international experts. We updated searches in all databases in March 2009, January 2013, February 2014 and June 2016.

The Cochrane Highly Sensitive Search Strategy for identifying randomised studies in MEDLINE, sensitivity-maximising version, Ovid format (Higgins 2008) was combined with the search terms in the Appendices to identify randomised controlled studies in MEDLINE. The MEDLINE search strategy was adapted for use in the other databases searched. The search strategies are documented in Appendix 1; Appendix 2 and Appendix 3. We did not confine our search to English language publications.

In addition, we searched the following databases of ongoing studies: <http://www.controlled-trials.com>; <http://www.clinicaltrials.nci.nih.gov>; <http://www.eortc.be>; <http://www.update-software.com/National/nrr-frame.html> and <http://www.CenterWatch.com>.

Searching other resources

We handsearched reference lists from studies selected by electronic searching to identify further relevant studies. We also handsearched published abstracts from conference proceedings from the European Society for Medical Oncology from 1978 (published in the *Annals of Oncology*), the European Council of Clinical Oncology from 1981 (published in the *European Journal of Cancer*), as well as the American Society for Clinical Oncology from 1981. All searches were updated in June 2016.

Data collection and analysis

Selection of studies

Two independent review authors initially scanned the title, abstract section, and keywords of every record retrieved. We retrieved full-text articles for further assessment if the information given suggested that the study included participants with histologically confirmed, inoperable adenocarcinoma of the stomach or gastroesophageal junction, used random allocation to the comparison groups and compared the following.

1. Best supportive care (BSC) versus chemotherapy plus BSC.
2. Combination versus single-agent chemotherapy.
3. 5-FU/cisplatin/anthracycline-containing combinations versus 5-FU/cisplatin combinations (without anthracyclines).
4. 5-FU/cisplatin/anthracycline-containing combinations versus 5-FU/anthracycline combinations (without cisplatin).
5. Irinotecan versus non-irinotecan-containing regimens.
6. Docetaxel versus non-docetaxel-containing regimens.
7. Capecitabine versus 5-FU-containing regimens.
8. Oxaliplatin versus the same regimen including cisplatin.
9. Taxane-platinum-fluoropyrimidin combinations versus taxane-platinum (without fluoropyrimidine).
10. S-1 versus 5-FU-containing regimens.

If there was any doubt regarding these criteria from the information given in the title and abstract, we retrieved the full-text article for clarification. If differences in opinion existed, they were resolved by discussion.

Data extraction and management

Two review authors independently extracted details of study population, interventions, and outcomes. We resolved differences in data extraction by consensus with a third review author, referring

back to the original article. If data were missing in a published report, we contacted the primary author.

Data extraction included the following items.

1. General information: title, authors, source, contact address, country, published/unpublished, language and year of publication, sponsoring of study.
2. Study characteristics, including design, duration/follow up, and quality assessment criteria as specified above.
3. Participants: inclusion and exclusion criteria, sample size, baseline characteristics, similarity of groups at baseline, withdrawals, and losses to follow-up.
4. Interventions: dose, route, timing of chemotherapy, and comparison intervention.
5. Outcomes: hazard ratios and their 95% confidence intervals or standard error, log rank χ^2 , log rank P values, number of events, number of participants per group, median, one-, two- and three-year survival rates and five-year survival rates in participants with locally advanced, secondary resectable tumours.

Assessment of risk of bias in included studies

In this updated version of the review, we independently assessed the risk of bias of the included studies using the 'Risk of bias' assessment tool described in Chapter 8 of the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011). We compared the evaluations, and discussed and resolved any inconsistencies between the review authors' decisions.

We rated the following domains separately for each of the included studies as 'low risk of bias', 'high risk of bias', and 'unclear' when the risk of bias was uncertain or unknown:

1. generation of allocation sequence ('sequence generation');
2. concealment of allocation ('allocation concealment');
3. prevention of knowledge of the allocated interventions during the study ('blinding');
4. methods used to address incomplete outcome data;
5. selective outcome reporting;
6. other sources of bias that could put a study at high risk of bias, including whether a calculation of sample size was carried out including baseline comparability. We considered tumour stage (advanced versus metastatic disease) and activity index (Eastern Cooperative Oncology Group status 0 to 1 versus 2 to 3), as well as the number of organs involved in metastatic disease (one versus more than one) as the most important prognostic factors. We considered a difference of more than 15% between study arms as an important difference. For age, as a further important factor, we considered baseline differences of five years as important. We also assessed intention-to-treat (ITT) analysis. We considered ITT analysis as randomised analysis, with the analysis restricted to participants who received at least one cycle of chemotherapy, and for which survival data were available. Alternatively, we also considered studies including all participants as randomised in the analysis as ITT. In addition, we analysed the risk of bias and described this in the 'Risk of bias' tables.

These assessments are reported in the 'Risk of bias' table for each individual study in the '[Characteristics of included studies](#)' section

of the review, and in the '[Risk of bias in included studies](#)' section of this review.

Measures of treatment effect

Data analysis

We estimated hazard ratios (HR) and 95% confidence intervals (CI) as relevant effect measures directly or indirectly from the given data (Altman 2001). For each individual study, we extracted HRs and their variances. If the figures were not given directly, methods of indirect determination were used. HRs can be estimated (under some assumptions) from log rank χ^2 , from log rank P values, from observed to expected event ratios and from ratios of median survival times or time point survival rates (Machin 1997; Parmar 1998; Tierney 2007). In several instances, medians and/or number of events had to be read from the graphs. If both medians and survival rates at fixed time points were given, the medians were preferred. If we had to pool several arms of a study, we approximated the common median by the weighted mean of the medians given for the various arms.

For instance, statistical measures were taken to avoid double-counting the irinotecan-treated population in Bouche 2004, which was a three-arm study which compared 5-fluorouracil *versus* 5-fluorouracil plus cisplatin *versus* 5-fluorouracil plus irinotecan. For Comparison 5, we estimated the hazard ratios for the irinotecan-containing arm (N = 45) versus the non-irinotecan-containing arms (N = 89), which was not provided in the original report by Bouche 2004.

Unit of analysis issues

Participants were individually randomised into two or more treatment groups. The effect of the intervention was measured and analysed on the basis of single measurements for each outcome for each participant. For studies with more than one intervention arm, we combined groups to create a single pair-wise comparison as follows: the single-agent therapy arm with the pooled results of both combination chemotherapy arms; the combination-therapy arm was compared with the pooled results of both single agent arms; or the pooled results of the sequential therapy arms were compared with the pooled results of the concurrent therapy arms.

Dealing with missing data

We attempted to contact investigators to obtain missing data.

Assessment of heterogeneity

We assessed the heterogeneity in each pair-wise comparison by assessing the Higgins I^2 (Higgins 2003), the χ^2 test with significance set at a P value less than 0.1, and by visual inspection.

Data synthesis

We used the fixed-effect model for meta-analysis, with overall survival as the primary outcome measure. Where heterogeneity levels were high ($I^2 > 20\%$ or P value < 0.1), we used a random-effects model (see [Differences between protocol and review](#)). We used Review Manager software for data synthesis (RevMan). . Previously SAS was used for more sophisticated analysis. We recalculated (or at least approximated) all outcomes concerning overall survival by using hazard ratios.

Quality of Evidence (GRADE) and 'Summary of findings' tables

We used the GRADEprofiler (GRADEpro) software to assist with the preparation of the 'Summary of findings' tables. The 'Summary of findings' tables provide key information about the pooled estimate of the magnitude of the effect in relative terms, assumed and control risks, numbers of participants and studies addressing each important outcome, and the quality of evidence for the comparisons for each outcome.

We included the following outcomes in the 'Summary of findings' tables.

1. Overall survival
2. Tumour response
3. Progression-free survival; and/or time-to-progression; and/or time-to-treatment failure
4. Treatment-related death
5. Treatment discontinuation due to toxicity

We assessed the quality of evidence as 'High', 'Moderate', 'Low' or 'Very Low' using the Grades of Recommendation, Assessment, Development and Evaluation (GRADE) methodology, which evaluates the totality of included studies for their risk of bias (study limitations), consistency, imprecision, indirectness, and publication bias.

Subgroup analysis and investigation of heterogeneity

In seeking statistical heterogeneity between studies, we performed Cochrane's Q-test (with a significance threshold of $\alpha = 0.1$). Additionally, we calculated heterogeneity quantitatively

(Thompson 2002). We considered the following factors as possible sources of heterogeneity:

1. differences in prognostic factors;
2. quality of studies;
3. second-line therapy permitted versus no second-line therapy;
4. Asian versus non-Asian studies;
5. substitutive, additive, and other comparisons in comparisons 5 and 6.

Sensitivity analysis

We conducted a sensitivity analysis by repeating the primary analysis and investigated the influence of risk of bias, adequate allocation concealment, excluding those studies which were conducted in Asia and studies with second-line therapies.

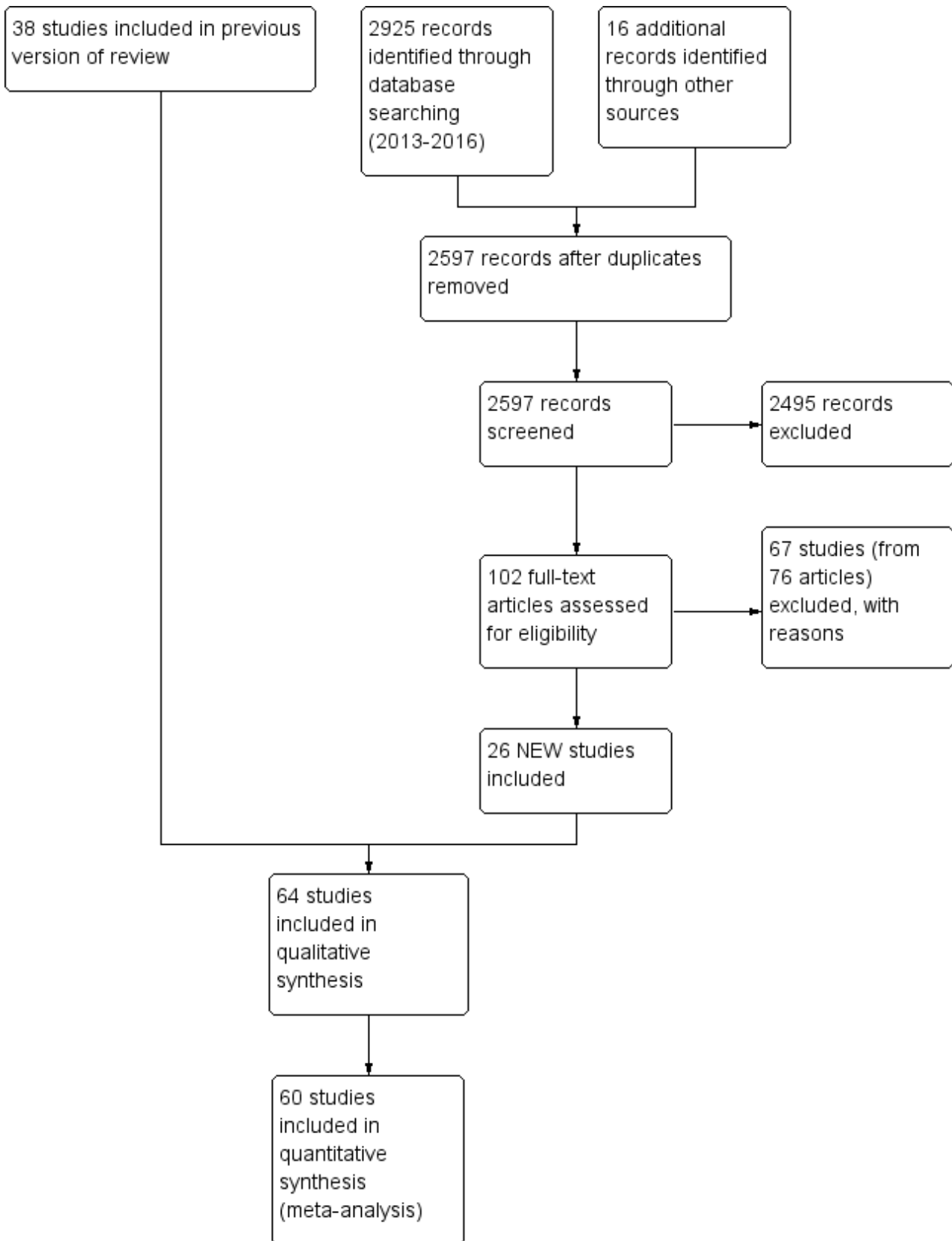
RESULTS

Description of studies

Results of the search

We identified a total of 2925 records through electronic searches of CENTRAL, MEDLINE, Embase and databases of clinical trials. After removing duplicates, 2597 records remained. We excluded 2495 references which were clearly irrelevant through screening titles and reading abstracts. We retrieved 102 references for further assessment. We excluded 195 studies and are listed in the table [Characteristics of excluded studies](#). Twenty-six new studies were identified for inclusion. Please see [Figure 1](#) for the flowchart of the systematic search performed in June 2016.

Figure 1. Study flow diagram: review update



Included studies

Three eligible studies with 184 participants were identified for analysis of **comparison 1**: chemotherapy versus best supportive care (BSC). The study by [Scheithauer 1996](#) was published as an abstract only, but all relevant information was provided by the author. For more details of the included studies, please see [Characteristics of included studies](#).

Approximately 50% of the study investigators provided further information. Data about survival and response rates were given in most publications. Information about second-line therapy was either reported in the text or provided by the authors only in a limited number of studies.

Twenty-three studies, which included 4447 participants, were included in the analysis of **comparison 2**, which is combination versus single-agent chemotherapy. In studies that included more than one single or combination chemotherapy arm, different arms were combined in the analysis as specified below. **Comparisons 3 and 4** included 579 and 1147 participants in four and seven studies. **Comparison 5** included 10 studies with a total of 2135 participants and **comparison 6** includes at present overall survival data from eight studies including a total of 2001 participants. **Comparisons 7 and 8** included 732 and 1105 participants from two and five studies, respectively. Two new **comparisons (9) and (10)** with 482 and 1793 participants from three and four randomised studies, respectively were added in the current update.

It should be noted that some studies may appear in more than one comparison if they meet relevant criteria for inclusion. For instance, [Hironaka 2016](#); [Koizumi 2014](#); [Komatsu 2011](#); [Narahara 2011](#); [Ochendusko 2015](#) were included in two comparisons; while [Boku 2009](#) was included in three comparisons.

Participants

The median age of the participants in the population of studies included in the analysis of **comparisons 1 and 2** was in the range of 56 to 67 years. The proportion of participants with metastatic disease was between 62% ([Cullinan 1985](#)) and 100% ([Bouche 2004](#); [Koizumi 2008](#); [Yamamura 1998](#)). When comparing the different arms of one study, a difference in the proportion of participants with advanced versus metastatic disease greater than 15% between study arms was identified in only one study ([Popov 2002](#)), with a larger number of metastatic participants in the combination chemotherapy arm (90% versus 73%). Performance status was well-balanced in all studies with no differences greater than 15% between study arms. The percentage of participants with ECOG 2+3 was in the range of 0% to 48%. Thirteen studies, which included 3182 participants ([Boku 2009](#); [Hironaka 2016](#); [Koizumi 2008](#); [Koizumi 2014](#); [Komatsu 2011](#); [Lu 2014](#); [Narahara 2011](#); [Nishikawa 2012](#); [Ohtsu 2003](#); [Shirao 2013](#); [Wang 2013](#); [Wu 2015](#); [Yamamura 1998](#)), were conducted in Asia.

Regarding **comparisons 3 and 4**, the median age of participants included in these 11 studies was between 58 and 65 years. Between 46% ([Kikuchi 1990](#)) and 90% ([Cascinu 2011](#); [Kim 2001](#)) of participants had metastatic disease, the percentage of participants with ECOG 2+3 was between 6% ([Cascinu 2011](#)) and 88% ([Kikuchi 1990](#)). The percentage of participants with advanced versus metastatic disease was well-balanced in all studies included in these two comparisons.

The participants in **comparison 5** had a median age between 58 ([Dank 2008](#)) and 70 ([Komatsu 2011](#)) years in the different study arms, with the majority of participants having metastatic disease.

The median age of the participants in **comparison 6** was between 55 ([Roth 2007](#); [Van Cutsem 2006](#)) and 70 years ([Al-Batran 2013](#)). The percentage of participants with metastatic disease was between 69% ([Al-Batran 2013](#); [Ochendusko 2015](#)) and 98% ([Thuss-Patience 2005](#)), with the largest study having 97% of participants with metastatic disease and a median age of 55 years ([Van Cutsem 2006](#)). Most participants in these studies had a performance status of 0 or 1.

Regarding **comparisons 7 and 8**, the median age of participants included in these studies was between 55 ([Ocvirk 2012](#)) and 65 years ([Yamada 2015](#)). Between 62% ([Kim 2014](#)) and 100% ([Li 2016](#)) had metastatic disease and most participants had a performance status of 0 or 1. Only in [Popov 2008](#) the percentage of participants with ECOG 2-3 was 29%.

The median age of the participants included in **comparisons 9 and 10** was between 54 ([Ajani 2005](#); [Huang 2013](#); [Li 2015](#)) and 76 years ([Boku 2009](#)), between 86% ([Roth 2007](#)) and 95% to 100% of participants ([Ajani 2005](#); [Ajani 2010](#); [Chen 2015](#)) had metastatic disease. Most participants in these studies had a performance status of 0 or 1.

Groups of participants were well-balanced regarding the most important prognostic factors as specified above in all studies included into **comparisons 3 to 8**.

Interventions

Participants were individually randomised into two or more treatment groups. The effect of the intervention was measured and analysed on the basis of single measurements for each outcome for each participant. In 11 studies, more than two groups with different interventions were compared ([Boku 2009](#); [Bouche 2004](#); [Cullinan 1985](#); [Cullinan 1994](#); [Hironaka 2016](#); [Loehrer 1994](#); [Lutz 2007](#); [Nishikawa 2012](#); [Ohtsu 2003](#); [Roth 2007](#)).

In the studies by [Bouche 2004](#), [Lutz 2007](#), [Ohtsu 2003](#), and [Hironaka 2016](#) the single-agent therapy arm was compared with the pooled results of both combination chemotherapy arms. In [Loehrer 1994](#), [Cullinan 1994](#), and [Boku 2009](#), the combination-therapy arm was compared with the pooled results of both single-agent arms. In [Nishikawa 2012](#), the pooled results of the sequential therapy arms were compared with the pooled results of the concurrent therapy arms. All these studies were included in the comparison of single-agent versus combination chemotherapy.

Furthermore, the irinotecan-containing combination chemotherapy was compared to the pooled results of the non-irinotecan-containing combination chemotherapies and included in comparison **5** in 10 studies ([Boku 2009](#); [Bouche 2004](#); [Dank 2008](#); [Komatsu 2011](#); [Li 2016](#); [Moehler 2005](#); [Moehler 2010](#); [Narahara 2011](#); [Roy 2012](#); [Sugimoto 2014](#)), and the pooled results of the docetaxel-containing chemotherapies were compared with the non-docetaxel-containing combination chemotherapy and included in comparison **6** in eight studies ([Al-Batran 2013](#); [Koizumi 2014](#); [Ochendusko 2015](#); [Ridwelski 2008](#); [Roth 2007](#); [Thuss-Patience 2005](#); [Van Cutsem 2006](#); [Wang 2016](#)).

All three studies included in the analysis of **comparison 1** used combination therapy regimens in the chemotherapy arm (Murad 1993: FAMTX; Pyrhönen 1995: FEMTX; Scheithauer 1996: 5-FU/LV/ Epirubicin).

Regarding **comparison 2**, most studies used 5-FU in the single-agent arm. In six of 17 studies (Barone 1998; Colucci 1995; Cullinan 1985; Cullinan 1994; De Lisi 1986; Loehrer 1994), 5-FU was given as a bolus in doses of approximately 500 mg/m² days one to five every four weeks. A continuous infusion regimen was used in two studies, with either 2600 mg/m² every two weeks (Popov 2002), 800 mg/m² per day (Ohtsu 2003) on days one to three every four weeks or a bolus of 400 mg/m² 5-FU, followed by 600 mg/m² as a two-hour continuous infusion on days one and two every two weeks (Bouche 2004). One study (Levi 1986), applied doxorubicin 60 mg/m² every four weeks in the single-agent arm. In the study by Loehrer 1994, the results from two single-agent arms (5-FU bolus 500 mg/m² days one to five and epirubicin 90 mg/m² day one every four weeks) were combined in the analysis. In addition, in the studies by Bouche 2004, Lutz 2007, Nishikawa 2012, and Ohtsu 2003, the results of two combination therapy arms (LV5FU2/cisplatin and LV5FU2/irinotecan, D-FU/FA and HD-FU/FA/cisplatin, 5-FU/paclitaxel and S-1/paclitaxel, 5-FU/cisplatin, and tegafur/mitomycin C) were combined in the analysis. Nishikawa 2012 used 5-FU in doses of 800 mg/m² in days one to five every four weeks in one group or daily S-1 in doses of 80 mg/m² for four weeks and a two-week rest in the single-agent arm until progression. This therapy was followed by paclitaxel (80 mg/m² on days one, eight and 15 every four weeks). A similar regimen of S-1 was used in Narahara 2011 and Komatsu 2011. Boku 2009 used the same 5-FU regimen in one arm or lower doses of S-1 (40 mg/m² for four weeks and a two-week rest) in the single-agent arm. Results of two arms were combined in the single-agent arm in Boku 2009; Cullinan 1994 and Nishikawa 2012. Wang 2013 used S-1 according to body-surface area with 40 mg twice daily for participants with a body surface area greater than 1.25 and lower than 1.5 for two out of four weeks. Koizumi 2008 and Hironaka 2016 as well used the oral prodrug S-1.

In six of 23 studies, combination chemotherapy arms did contain an anthracycline (epirubicin 60 mg/m² every three weeks or 90 mg/m² every four weeks or doxorubicin 40 mg/m² every four to seven weeks) in addition to 5-FU. Non 5-FU-based combination chemotherapy regimens, which instead included etoposide, irinotecan and cisplatin, S-1 and irinotecan, S-1 and paclitaxel or an anthracycline and cisplatin were applied in six studies (Barone 1998; Boku 2009; Komatsu 2011; Narahara 2011; Popov 2002; Wang 2013).

In **comparisons 3** and **4**, regimens containing 5-FU, an anthracycline and cisplatin were mainly FAP (5-FU bolus 300 mg/m² either days one, eight, 15, 22 or days one to five, adriamycin 25 to 40 mg/m² and cisplatin 60 to 100 mg/m² once every three to five weeks) (Cullinan 1994; GITSG 1988; Kikuchi 1990), and PELF (cisplatin 40 mg/m² days one and five, epirubicin 30 mg/m² days one and five, leucovorin 200 mg/m² and 5-FU bolus 300 mg/m² days one to four every eight weeks) (Cocconi 1994) in studies published before 1995 (Kim 2001; Ross 2002; Webb 1997). used mostly ECF (epirubicin 50 mg/m² and cisplatin 60 mg/m² once every three weeks, with 5-FU as a continuous intravenous infusion of 200 mg/m² for up to six months) or LdCF (pegylated liposomal doxorubicin 20 mg/m² and cisplatin 50 mg/m² once every two weeks, with 5-

FU 400 mg/m² bolus followed by 600 mg/m² as 22-hour continuous infusion on days one and two every two weeks) in Cascinu 2011.

In **comparison 5**, irinotecan was given in the following studies to substitute either cisplatin (Bouche 2004; Dank 2008; Moehler 2010), etoposide (Moehler 2005) or 5-FU (Roy 2012) or paclitaxel (Sugimoto 2014). In contrast, irinotecan was given in addition to the treatment in the other arm in studies by Bouche 2004; Komatsu 2011; and Narahara 2011; or as other comparisons (Li 2016). Irinotecan was given weekly at 80 mg/m² for six weeks every 50 days in combination with infusional 5-FU 2000 mg/m² and FA 500 mg/m² (Dank 2008; Moehler 2005). Moehler 2010 used irinotecan 250 mg/m² on day one in combination with capecitabine 1000 mg/m² orally days one to 14 every 22 days and Roy 2012 used irinotecan 250 mg/m² on day one in combination with docetaxel 60 mg/m² as infusion on day one every 22 days. Bouche 2004 used irinotecan at 180 mg/m² in combination with infusional 5-FU every two weeks. Komatsu 2011 and Narahara 2011 used doses of 75 mg/m² and 80 mg/m² on days one and 15 every four weeks or six weeks in combination with oral S-1 (initial doses of 40 mg/m² to 60 mg/m² twice daily on days one to 14 every four weeks or 80 mg/m²/day on days one to 21 every six weeks). In subsequent cycles, doses were varied according to the most severe adverse events during the preceding cycle.

In **comparison 6**, docetaxel was given to substitute either epirubicin and cisplatin (Roth 2007; Thuss-Patience 2005) or 5-FU and leucovorin (Ridwelski 2008). On the other hand, it was given in addition to the treatment in the other study arm in the studies by Wang 2016, Koizumi 2014, Van Cutsem 2006 and Al-Batran 2013. The largest studies included in comparison 6's meta-analysis of overall survival used the three-drug regimen DCF (docetaxel 75 mg/m² intravenously day one, cisplatin 75 mg/m² intravenously day one, 5-FU 750 mg/m² as a 24-hour infusion) on days one to five every three weeks (Van Cutsem 2006), and the two-drug regimen of docetaxel 75 mg/m² intravenously day one, in combination with cisplatin 75 mg/m² intravenously on day one every three weeks (Ridwelski 2008). In Koizumi 2014, docetaxel (40mg/m² intravenously on day one) was given with S-1 (tailored to body surface area; days one to 14) every 21 days. In Roth 2007, the DCF regimen was used as described previously. Sadighi 2006 and colleagues used a modification of DCF with reduced doses of docetaxel and cisplatin (both at 60 mg/m²) every three weeks. Thuss-Patience 2005 applied docetaxel 75 mg/mg/m² intravenously on day one in combination with 5-FU 200 mg/m²/day over 24 hours on days one to 21 every three weeks. Al-Batran 2013 used docetaxel 50 mg/m² intravenously on day one in combination with oxaliplatin 85 mg/m² and leucovorin 200 mg/m² followed by 5-FU 2600 mg/m² as a 24-hour continuous infusion.

Five studies (Kang 2009; Li 2016; Ocvirk 2012; Ochendusko 2015; Van Cutsem 2015) are eligible for **comparison 7**. In Kang 2009 and Ocvirk 2012, the oral 5-FU prodrug capecitabine (1000 mg/m² or 825 mg/m² twice daily on days one to 14 of a 21-day regimen) was compared with 5-FU (both in combination with cisplatin). In **comparison 8**, oxaliplatin was given once at 85 mg/m² in two weeks and compared with cisplatin 50 mg/m² in Al Batran 2008 and Popov 2008. Both agents were combined with FU/ leucovorin in Al Batran 2008 and Popov 2008. In Kim 2014 a combination of weekly docetaxel (35mg/m²) on days one and eight every three weeks, in combination with either cisplatin (60 mg/m²) or oxaliplatin (120 mg/m²) on day one was administered. In Yamada 2015, S-1 (twice

daily for the first three weeks of a five-week cycle) plus (cisplatin 60 mg/m² on day eight) was compared against S-1 (twice daily for first two weeks of a three-week cycle) plus oxaliplatin (100 mg/m² infused for two hours on day one). For **comparison 9**, three studies (Ajani 2005; Roth 2007; Van Cutsem 2015) are available. Fluorouracil 200 mg/m²/day was given as a 24-hour continuous infusion in or in doses of 750 mg/m²/day on days one to five every three weeks in Roth 2007. It was combined with docetaxel and cisplatin in Ajani 2005 and Roth 2007, and docetaxel and oxaliplatin in Van Cutsem 2015.

In **comparison 10**, S-1 50 mg/m² was given orally in two daily doses on days one to 21 of a four-week cycle (Ajani 2010), or in a lower dosage of 40 mg/m² orally in two daily doses on days one to 28 of a six-week cycle (Boku 2009) and compared with continuous infusions of 5-FU in doses of 1000 mg/m²/24 hours as 120-hour infusion (Ajani 2010) or 800 mg/m²/day on days one to five, respectively. Of note, S-1 was combined with 75 mg/m² of cisplatin every three weeks, and 5-FU with 100 mg/m² of cisplatin every three weeks in the study by Ajani 2010. Huang 2013 compared the combination of weekly paclitaxel (60 mg/m²) on days one, eight, and 15 every four-week cycle and S-1 (80 mg/m² to 120 mg/m²), dependent on the body-surface area for two out of four weeks, with the same regimen of paclitaxel, but with 5-FU (500 mg/m², in combination with leucovorin 20 mg/m² on days one to five every four weeks).

Outcomes

Median survival and response rates were the outcomes most commonly described in the included studies. The newer studies reported progression-free survival instead of time to progression. Toxicity was not always classified according to WHO or NCI-CTC and was reported in different ways (per number of participants, per number of cycles and only the worst toxicity per participant). A comparison across studies was therefore not possible. For this reason, the numbers of participants who discontinued treatment due to toxicity as well as the numbers of treatment-related deaths were analysed. Information about second-line therapies and secondary resectability was unavailable for most studies. Details are listed in the [Characteristics of included studies](#) table.

Excluded studies

Please see [Characteristics of excluded studies](#). Reasons for exclusion of references in the updated search are specified in [Figure 1](#) according to the recommendations of the PRISMA statement (Moher 2009).

According to the protocol, we excluded studies in which cross-over after failure was encouraged or planned. Information about

second-line therapy was generally unavailable in most first-line studies. However, in some large recent studies full information about second-line therapy was provided after contacting the first author and a limited number of participants did in fact cross over in the second-line therapy. After discussion and balancing the loss of information when excluding these studies against the possible bias caused by a limited number of participants crossing over, we decided to include studies in which the number of participants who crossed over between study arms was less than 10% of the whole study population. Exclusion of these studies would have provoked a bias in favour of studies where less information was available. Two studies which have repeatedly been quoted in the literature are especially mentioned: Glimelius 1994 was excluded because of cross-over, as the research ethics committee had requested that chemotherapy had to be provided to participants upon request in the BSC group, and 12 of 30 participants in the BSC group finally received chemotherapy. Kim 1993 was excluded since the allocation of participants was done by alternate allocation (information provided by author YSP). Several studies currently published as abstracts only were not included at this stage, because full information and final results were currently unavailable or were not provided after contacting the author or sponsor. They are classified as 'Studies awaiting classification'. We excluded studies using oral 5-FU because of its varying bioavailability and unreliable effect. The landmark 'REAL-2' study (Cunningham 2008), which evaluated the non-inferiority of oxaliplatin as compared to cisplatin and of capecitabine as compared to 5-FU, was not included in this meta-analysis after discussion because it included participants with squamous cell cancer of the oesophagus (more than 10%), which were not eligible according to the inclusion criteria for this meta-analysis. We sought separate data on participants with adenocarcinoma only, but they were not provided by the study investigator. The study by Sadighi 2006 could not be included in the meta-analysis of overall survival because published data for calculation of the hazard ratio is not sufficient at present, but data on the other outcomes were included. The study by Park 2006 was not included in the comparison of docetaxel versus non-docetaxel-containing regimens because both study arms included a taxane. Thus, the analyses are essentially a comparison of docetaxel and paclitaxel. If further studies relevant for this comparison are published in the future, a separate comparison of paclitaxel versus docetaxel-containing regimens will be included in the meta-analysis. The studies Gubanski 2010 and Guimbaud 2014 were excluded because of systematic cross-over between study arms.

Risk of bias in included studies

We summarised the overall risk of bias in [Figure 2](#) and [Figure 3](#).

Figure 2. 'Risk of bias' summary: review authors' judgements about each risk of bias item for each included study.

	Random sequence generation (selection bias)	Allocation concealment (selection bias)	Incomplete outcome data (attrition bias): efficacy	Incomplete outcome data (attrition bias): safety	Selective reporting (reporting bias)	Other bias	Blinded review of CT/MRI-scans?
Ajani 2005	+	+	+	+	?	?	+
Ajani 2010	+	+	+	+	+	+	+
Al Batran 2008	?	?	+	+	?	-	?
Al-Batran 2013	+	+	+	+	+	+	-
Barone 1998	+	+	+	+	+	-	-
Boku 2009	+	+	+	+	+	+	+
Bouche 2004	?	+	+	+	+	?	?
Cascinu 2011	+	+	+	+	+	+	+
Chen 2015	+	?	?	?	?	?	?
Cocconi 1994	?	+	+	+	+	+	+
Cocconi 2003	?	+	?	?	?	?	?
Colucci 1995	?	?	+	+	+	?	-
Cullinan 1985	?	?	+	+	+	-	-
Cullinan 1994	?	?	+	+	?	-	?
Dank 2008	+	?	+	+	+	-	+
De Lisi 1986	?	+	+	+	+	?	-
Dong 2014	+	?	?	?	-	?	?
GITSG 1988	?	?	?	?	?	?	?
Hironaka 2016	+	+	+	+	+	?	+
Huang 2013	?	+	+	+	+	?	?
Kang 2009	+	?	+	+	+	+	+
Kikuchi 1990	?	+	+	+	+	?	-

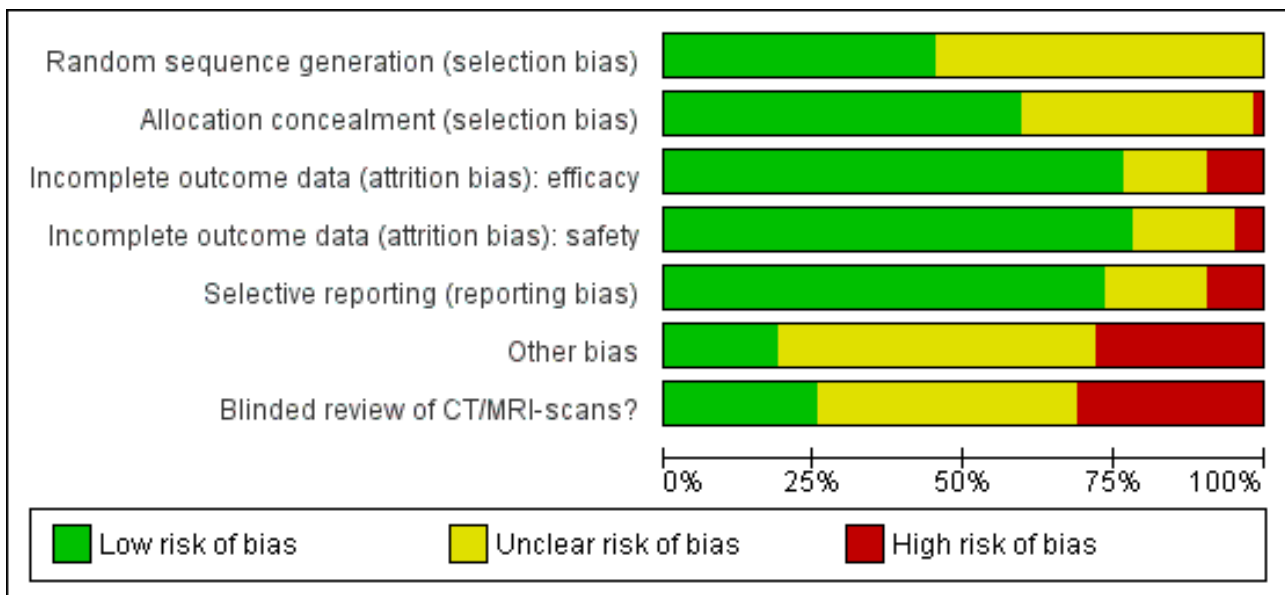
Figure 2. (Continued)

Kikuchi 1990	?	+	+	+	+	?	-
Kim 2001	?	?	?	?	+	?	-
Kim 2014	?	?	+	+	+	+	?
Koizumi 2008	+	+	+	+	+	+	+
Koizumi 2014	?	?	+	+	+	-	?
Komatsu 2011	+	?	?	+	+	-	-
KRGGC 1992	?	+	-	?	+	?	?
Levi 1986	?	+	-	-	+	?	?
Li 2014	+	?	-	-	?	?	?
Li 2015	?	-	?	?	?	-	-
Li 2016	?	?	?	?	?	?	?
Loehrer 1994	+	+	-	?	-	+	-
Lu 2014	+	+	+	+	?	?	?
Lutz 2007	+	+	+	+	+	-	-
Moehler 2005	?	+	+	+	+	+	?
Moehler 2010	?	+	+	+	+	-	?
Murad 1993	?	?	-	?	+	?	?
Narahara 2011	+	+	+	?	+	-	?
Nishikawa 2012	+	+	+	+	+	?	?
Ochendusko 2015	?	?	+	+	?	-	?
Ocvirk 2012	?	+	+	+	+	-	-
Ohtsu 2003	+	+	+	+	+	+	?
Popov 2002	+	?	+	+	+	?	+
Popov 2008	?	?	-	+	+	?	+
Pyrhönen 1995	+	+	+	+	+	-	-
Ridwelski 2008	+	+	+	+	+	?	-
Ross 2002	?	?	+	+	+	?	-
Roth 1999	?	?	+	+	+	?	-
Roth 2007	?	+	+	+	+	?	+
Roy 2012	?	?	+	+	+	?	?
Sadighi 2006	?	?	?	-	-	?	?

Figure 2. (Continued)

	+	+	-	-	+	+
Sadighi 2006	?	?	?	-	-	?
Scheithauer 1996	+	+	+	+	+	-
Shirao 2013	+	+	+	+	-	?
Sugimoto 2014	?	?	+	+	+	+
Thuss-Patience 2005	+	+	+	+	+	+
Van Cutsem 2006	?	+	+	+	+	+
Van Cutsem 2015	?	+	+	+	+	?
Wang 2013	+	+	+	+	+	-
Wang 2016	?	+	+	+	+	-
Webb 1997	?	?	+	+	+	-
Wu 2015	+	+	+	+	-	?
Yamada 2015	+	+	+	+	+	+
Yamamura 1998	?	+	+	+	-	?

Figure 3. 'Risk of bias' graph: review authors' judgements about each risk of bias item presented as percentages across all included studies.



Allocation

The risk of selection bias due to issues with random sequence generation was not stated in most instances (n = 35/64, 55%), while the remaining studies which described the allocation sequence generation approach used acceptable, unbiased methods (n = 29/64, 45%).

Blinding

Potential bias arising from allocation concealment was low in 38 studies (59%), unclear in 25 studies (39%), and high in one study (2%).

Incomplete outcome data

The risk of bias due to incomplete efficacy data was low in 49 studies (77%), unclear in nine studies (14%), and high in six studies (9%). On the other hand, incomplete safety concern was a low concern in 50 studies (78%), unclear in 11 studies (17%), and high in three studies (5%).

Selective reporting

The potential for selective reporting was deemed to be low in 47 studies (73%), unclear in 11 studies (17%), and high in six studies (9%).

Other potential sources of bias

The risk of bias due to lack of blinded or independent radiological review was a low concern in 16 studies (25%), unclear in 28 studies (44%), and high in 20 studies (31%). The potential for bias due to other causes was assessed to be low in 12 studies (19%), unclear in 34 studies (53%), and high in 18 studies (28%).

Effects of interventions

See: [Summary of findings for the main comparison](#) Chemotherapy versus best supportive care for advanced gastric cancer; [Summary of findings 2](#) Combination versus single-agent chemotherapy for advanced gastric cancer; [Summary of findings 3](#) 5-FU/cisplatin/anthracycline combinations versus 5-FU/cisplatin combinations (without anthracyclines) for advanced gastric cancer; [Summary of findings 4](#) 5-FU/cisplatin/anthracycline combinations versus 5-FU/anthracycline combinations (without cisplatin) for advanced gastric cancer; [Summary of findings 5](#) Irinotecan versus non-irinotecan-containing regimens for advanced gastric cancer; [Summary of findings 6](#) Docetaxel versus non-docetaxel-containing regimens for advanced gastric cancer; [Summary of findings 7](#) Capecitabine versus 5-FU-containing regimens for advanced gastric cancer; [Summary of findings 8](#) Oxaliplatin versus the same regimen including cisplatin for advanced gastric cancer; [Summary of findings 9](#) Taxane-platinum-fluoropyrimidine combinations versus taxane-platinum (without fluoropyrimidine) for advanced gastric cancer; [Summary of findings 10](#) S-1 versus 5-FU-containing regimens for advanced gastric cancer

(1) Chemotherapy versus best supportive care

Overall survival

A total of three studies (N = 184) reported overall survival ([Murad 1993](#); [Pyrhönen 1995](#); [Scheithauer 1996](#)). The overall hazard ratio (HR) of 0.37 (95% confidence interval (CI) 0.24 to 0.55, moderate-quality evidence) ([Analysis 1.1](#)) in favour of the chemotherapy arms demonstrates a convincing benefit over best supportive care (BSC) alone, which can be interpreted as an improvement in median survival from 4.3 months (weighted average in BSC) to 11 months (with chemotherapy). Cochrane's Q test (P = 0.19) as well as the index of homogeneity according to Thompson ($I^2 = 39.7$) demonstrate a significant statistical heterogeneity among the results of these studies, based on the differences in the chemotherapy regimens studied. A sensitivity analysis including only studies with adequate allocation concealment ([Pyrhönen 1995](#); [Scheithauer 1996](#)), does not change the overall HR of 0.37 (95% CI 0.19 to 0.70).

Secondary outcomes

Tumour response

Data were available for 88 participants in the chemotherapy arms of the three eligible studies. Response rates were between 33% ([Pyrhönen 1995](#)) and 50% ([Murad 1993](#)).

Time to progression

In the chemotherapy and BSC arms time to progression was 7.8 versus 2.7 (P = 0.0001) and 6.5 versus 2.0 (P = 0.0001) months in the studies by [Murad 1993](#) and [Scheithauer 1996](#) (N = 144). The overall HR was 0.31 (95% CI 0.22 to 0.43, moderate-quality evidence) ([Analysis 1.2](#)).

Secondary resectability

Information about secondary resectable participants was not given in the text or provided by the authors in any of the three studies.

Toxicity

In the study by [Murad 1993](#), WHO grade III/IV toxicities occurred in 22 of 128 cycles in 30 participants treated with chemotherapy, with one toxic death. [Pyrhönen 1995](#) described WHO gastrointestinal grade III/IV toxicities in 13 of 20 participants. Haematological toxicities occurred in the same frequency. [Scheithauer 1996](#) observed haematological grade III/IV toxicities in 12%, gastrointestinal grade III/IV toxicities in 21%, and other grade III/IV toxicities in 32.7% of 226 available cycles.

Quality of life

Quality of life was not analysed in any of the three included studies. [Pyrhönen 1995](#) assessed the palliative measures and observed an increased use of analgesics in the control versus treated participants after two months.

(2) Combination versus single-agent therapy

Overall survival

Twenty-three studies including 4447 participants were summarised in this meta-analysis ([Barone 1998](#); [Boku 2009](#); [Bouche 2004](#); [Colucci 1995](#); [Cullinan 1985](#); [Cullinan 1994](#); [De Lisi 1986](#); [Hironaka 2016](#); [Koizumi 2008](#); [Koizumi 2014](#); [Komatsu 2011](#); [Levi 1986](#); [Loehrer 1994](#); [Lu 2014](#); [Lutz 2007](#); [Narahara 2011](#); [Nishikawa 2012](#); [Ohtsu 2003](#); [Popov 2002](#); [Shirao 2013](#); [Wang 2013](#); [Wu 2015](#); [Yamamura 1998](#)). The overall HR in favour of combination chemotherapy (HR 0.84, 95% CI 0.79 to 0.89, moderate quality) provides evidence for a statistically significant, but modest survival benefit of combination versus single-agent chemotherapy in the studied regimens ([Analysis 2.1](#)). Cochrane's Q test for heterogeneity showed non-significant heterogeneity ($I^2 = 0\%$, P = 0.48), indicating that results of the different studies were consistent in their findings. As the chemotherapy regimens in studies published before 2000 might not have the same efficacy as modern regimens, the median survival difference between single and combination chemotherapy was calculated separately for studies published before the year 2000 and thereafter. For studies published before 2000, the weighted median survival was approximately 7.3 with combination therapy and 6.4 months with single-agent therapy. In studies published after 2000 ([Boku 2009](#); [Bouche 2004](#); [Hironaka 2016](#); [Koizumi 2008](#); [Koizumi 2014](#); [Komatsu 2011](#); [Lu 2014](#); [Lutz 2007](#); [Narahara 2011](#); [Nishikawa 2012](#); [Ohtsu 2003](#); [Popov 2002](#); [Wang 2013](#); [Wu 2015](#)), median survival was 11.6 months with

combination therapy, as compared to 10.5 months with single-agent therapy.

To evaluate the influence of second-line therapy, which was previously specified as a possible cause of heterogeneity, a second analysis was performed excluding the studies by [Boku 2009](#), [Koizumi 2008](#), [Koizumi 2014](#), [Narahara 2011](#), [Ohtsu 2003](#), and [Wang 2013](#) in which more than 50% of participants received a second-line therapy. Exclusion of these studies had no influence on heterogeneity ($I^2 = 0\%$ and $P = 0.56$) and the overall HR of 0.82 (95% CI 0.75 to 0.90) in favour of combination chemotherapy. Sensitivity analysis excluding those studies which were conducted in Asia ([Boku 2009](#); [Hironaka 2016](#); [Koizumi 2008](#); [Koizumi 2014](#); [Komatsu 2011](#); [Lu 2014](#); [Narahara 2011](#); [Nishikawa 2012](#); [Ohtsu 2003](#); [Shirao 2013](#); [Wang 2013](#); [Wu 2015](#); [Yamamura 1998](#)) resulted in a HR of 0.77 (95% CI 0.68 to 0.87) ($I^2 = 0\%$, $P = 0.57$), with no appreciable change in heterogeneity as compared to the original analysis. When only those studies with adequate allocation concealment were included in the analysis ([Barone 1998](#); [Boku 2009](#); [Bouche 2004](#); [De Lisi 1986](#); [Koizumi 2008](#); [Levi 1986](#); [Loehrer 1994](#); [Lu 2014](#); [Lutz 2007](#); [Narahara 2011](#); [Nishikawa 2012](#); [Ohtsu 2003](#); [Shirao 2013](#); [Wang 2013](#); [Wu 2015](#); [Yamamura 1998](#)), the resulting overall HR was 0.83 (95% CI 0.77 to 0.89) ($I^2 = 24\%$ and $P = 0.19$). For these reasons, the results of this comparison can be considered to be highly robust.

Secondary outcome measures

Tumour response

Data were available from 2833 participants in 18 eligible studies. The pooled objective response rate was 39% in the combination therapy arms versus 23% of the single-agent arms. The corresponding odds ratio (OR 2.30, 95% CI 1.94 to 2.72, high-quality evidence) confirms a statistically significant advantage in tumour response in favour of combination therapy ([Analysis 2.2](#)). Very low heterogeneity was observed across studies ($I^2 = 0\%$, $P = 0.60$).

Time to progression

Data from four studies with 720 participants were available. The overall HR for time to progression for combination versus single-agent chemotherapy was 0.69 (95% CI 0.55 to 0.87, moderate-quality evidence) ([Analysis 2.3](#)). Results across studies were consistent, with moderate heterogeneity ($I^2 = 23\%$, $P = 0.27$). Other studies ([Boku 2009](#); [Bouche 2004](#); [Hironaka 2016](#); [Koizumi 2014](#); [Lutz 2007](#); [Lu 2014](#); [Wu 2015](#)) reported progression-free survival instead of time to progression or time-to treatment failure ([Komatsu 2011](#); [Narahara 2011](#); [Nishikawa 2012](#)).

Secondary resectability

Only one study ([Colucci 1995](#)) reported on a single participant who became secondary resectable and was operated on with a pathologic complete remission.

Toxicity

Because of the different ways of reporting (per number of participants, per number of cycles or only the maximum toxicity per participant), grade I to IV toxicities can be compared only within, but not between studies. Overall, treatment-associated toxicities were higher in the combination chemotherapy arms, but this was usually not statistically significant. In contrast, the rate of treatment-associated deaths may be summarised across studies. Eighteen of 23 studies ($N = 3876$) in this comparison reported

treatment-related deaths ([Analysis 2.4](#)). The overall rate of toxic deaths in these studies was 1.1% versus 0.5% for combination versus single-agent therapy (OR 1.64, 95% CI 0.83 to 3.24). Six studies observed no treatment-related deaths ([Hironaka 2016](#); [Koizumi 2008](#); [Komatsu 2011](#); [Lu 2014](#); [Wang 2013](#); [Wu 2015](#)).

Quality of life

This was assessed in only one of these studies ([Bouche 2004](#)). All participants in the single-agent and both combination chemotherapy arms had a significant improvement in quality of life compared with pretreatment scores.

(3) 5-FU/cisplatin/anthracycline combinations versus 5-FU/cisplatin combinations (without anthracyclines)

Overall survival

This meta-analysis was based on 579 participants in four randomised studies ([Cascinu 2011](#); [Kim 2001](#); [KRGGC 1992](#); [Ross 2002](#)). The data from the largest study in this comparison (published by [Ross 2002](#)) which was included in this analysis were provided by the authors and include people with gastric and GE-junction adenocarcinoma only (the original publication included people with squamous cell carcinoma of the oesophagus as well). The resulting HR for overall survival of 0.74 (95% CI 0.61 to 0.89, moderate-quality evidence) demonstrates a statistically significant benefit with very low heterogeneity ($I^2 = 0\%$, $P = 0.63$) in overall survival in favour of the three-drug combination, with a weighted average survival of 9.9 and 8.6 months, respectively. Allocation concealment was adequate in all three studies included in this comparison and heterogeneity was non-significant ($I^2 = 0\%$; $P = 0.63$) ([Analysis 3.1](#)).

(4) 5-FU/cisplatin/anthracycline combinations versus 5-FU/anthracycline combinations (without cisplatin)

Summarising the results for the comparison of FU/cisplatin/anthracycline combinations versus FU/anthracycline (without cisplatin) results in a HR of 0.82 (95% CI 0.73 to 0.92, low-quality evidence) in favour of the three-drug regimen ([Analysis 4.1](#)). Combination chemotherapy arms only from the study by [Cullinan 1994](#) were included in this comparison. This meta-analysis, which included 1147 participants in seven studies ([Cocconi 1994](#); [Cocconi 2003](#); [Cullinan 1994](#); [GITSG 1988](#); [Kikuchi 1990](#); [Roth 1999](#); [Webb 1997](#)), once more confirms a overall survival benefit in favour of the three-drug combination, which corresponds to a difference in weighted mean average survival of approximately two months. A sensitivity analysis according to the quality score has only little impact on the resulting HR (0.79, 95% CI 0.68 to 0.91). There was only moderate heterogeneity ($I^2 = 28\%$, $P = 0.21$).

The two regimens containing FU, an anthracycline and cisplatin, which were evaluated in the largest number of participants are cisplatin, epirubicin, leucovorin, and FU administered as bolus (PELF; 184 participants) ([Cocconi 1994](#); [Cocconi 2003](#)) and epirubicin, cisplatin, and protracted venous-infusion FU (ECF; 327 participants) ([Kim 2001](#); [Ross 2002](#); [Webb 1997](#)). The rate of treatment-related deaths was 3.3% for PELF versus 0.6% for ECF (OR 5.36, 95% CI 1.1 to 27.4; Fisher's exact test, $P = .02834$), suggesting an increased toxicity of PELF. Quality of life was analysed in two studies evaluating ECF compared with FU, doxorubicin, and methotrexate ([Webb 1997](#)), as well as ECF compared with mitomycin, cisplatin, and FU ([Ross 2002](#)). Quality of life was superior in participants treated with ECF.

(5) Chemotherapy with irinotecan versus non-irinotecan-containing regimens

Overall survival

Ten studies (N = 2135) were summarised in this meta-analysis (Boku 2009; Bouche 2004; Dank 2008; Komatsu 2011; Li 2016; Moehler 2005; Moehler 2010; Narahara 2011; Roy 2012; Sugimoto 2014). To avoid double-counting the irinotecan-treated population in the study by Bouche 2004, we approximated a within-study hazard ratio of 0.72 (95% CI 0.30 to 1.72) for the study's irinotecan-treated population (N = 45) and non-irinotecan-treated population (N = 89). Overall, the pooled hazard ratio of irinotecan-containing regimens compared to non-irinotecan containing regimens was 0.87 (95% CI 0.80 to 0.95, high-quality evidence), with low heterogeneity ($I^2 = 0\%$, $P = 0.86$) (Analysis 5.1), and corresponds to pooled median survival times of 11.3 and 9.7 months, with a small, but significant benefit for the irinotecan-containing regimens. When only those studies with information about adequate allocation concealment are included in the analysis (Boku 2009; Bouche 2004; Moehler 2005; Moehler 2010; Narahara 2011; Roy 2012) the resulting overall HR was 0.84 (95% CI 0.76 to 0.93) ($P = 0.84$ and $I^2 = 0\%$) when all studies were pooled. For these reasons, the results of this comparison can be considered as robust. Further sensitivity analyses were not performed due to the low number of studies.

Next, the HR for overall survival was separately investigated for studies with substitutive (i.e. studies in which another chemotherapy was substituted by irinotecan), additive (i.e. studies in which irinotecan was added to other chemotherapies), and other comparisons of irinotecan and non-irinotecan-containing regimens (Analysis 5.1). The summary of the six studies (826 participants) with substitutive comparisons (Bouche 2004; Dank 2008; Moehler 2005; Moehler 2010; Roy 2012; Sugimoto 2014) results in a HR of 0.87 (95% CI 0.75 to 1.00, high-quality evidence), with low heterogeneity between study results ($I^2 = 0\%$, $P = 0.94$).

The meta-analysis of the three studies (Bouche 2004; Komatsu 2011; Narahara 2011), including a total of 500 participants, where irinotecan was given in addition to the treatment in the non-irinotecan-containing arm shows a non-significant benefit for participants treated with irinotecan (HR 0.88, 95% CI 0.76 to 1.03, low-quality evidence) and small heterogeneity between treatment effects of individual studies ($I^2 = 6\%$, $P = 0.34$). This result corresponds to a pooled median survival duration of 11.9 months with compared to 10.9 months without irinotecan. Bouche 2004 and Narahara 2011 stated a higher benefit of four and two months, respectively. However, Komatsu 2011 showed a disadvantage of three months for participants treated with irinotecan and S-1 compared to participants with S-1 monotherapy.

The meta-analysis of two studies (Boku 2009; Li 2016), which could neither be classified as substitutive nor additive, including a total of 809 participants, revealed a slight benefit for participants treated with irinotecan (HR 0.87, 95% CI 0.76 to 1.00, very low-quality evidence). However, the pooled result needs to be interpreted with caution in light of the considerable clinical differences between studies included under 'Other Comparisons' as well as their significant statistical heterogeneity ($I^2 = 65\%$, $P = 0.04$). For instance, Li 2016 allowed participants to switch to second-line therapy after failure of first-line. The study by Boku 2009 alone, which contributed the majority of participants (n = 704), showed a HR of 0.84, 95% CI 0.73 to 0.95 and a corresponding benefit in median

survival of around one month in favour of participants treated with irinotecan.

Secondary outcome measures

Tumour response

Data were available from 1266 participants in 10 eligible studies (Analysis 5.2). In six studies (756 participants) with substitutive comparisons (Bouche 2004; Dank 2008; Moehler 2005; Moehler 2010; Roy 2012; Sugimoto 2014) response rates were 38% and 30% (OR 1.53, 95% CI 0.93 to 2.50, low-quality evidence), with substantial heterogeneity ($I^2 = 56\%$, $P = 0.04$). In three studies (345 participants) with additive comparisons, pooled response rates of 38% and 22% were observed (OR 2.18, 95% CI 1.25 to 3.80, low-quality evidence), with moderate heterogeneity ($I^2 = 19\%$, $P = 0.29$). In two studies (165 participants) of other comparisons, response rates were 53% and 38% (OR 1.87, 95% CI 0.89 to 3.91, very low-quality evidence) with considerable heterogeneity ($I^2 = 26\%$, $P = 0.25$).

Progression-free survival

Most studies reported progression-free survival instead of time to progression. Again, a within-study hazard ratio was computed for the irinotecan-treated population (N = 45) and non-irinotecan-treated population (N = 89) in Bouche 2004 to avoid counting the irinotecan-treated population twice (HR = 0.59, 95% CI 0.50 to 0.68). Overall, the pooled hazard ratio of irinotecan-containing regimens compared to non-irinotecan-containing regimens was 0.76 (95% CI 0.69 to 0.84, high-quality evidence), with low heterogeneity ($I^2 = 0\%$, $P = 0.59$) (Analysis 5.3).

Among five studies (N = 741) with substitutive comparisons (Bouche 2004; Dank 2008; Moehler 2005; Moehler 2010; Sugimoto 2014), the hazard ratio for progression-free survival was 0.85 (95% CI 0.72 to 1.00, moderate-quality evidence) in favour of the participants treated with irinotecan (Analysis 5.3). The heterogeneity for this comparison was low ($I^2 = 0\%$, $P = 0.54$).

The single study (N = 90) from the additive comparisons, from which data for progression-free survival are available observed a large benefit (HR of 0.51, 95% CI 0.33 to 0.77, moderate-quality evidence) with a median PFS of 6.9 and 3.2 months for participants treated with and without irinotecan (Bouche 2004). A smaller difference in TTP was reported by Komatsu 2011, with a median TTP of 4.8 and 3.8 months for participants treated with and without irinotecan. Two additional studies belonging to other comparisons (Boku 2009; Li 2016) with a total of 809 participants stated a pooled benefit for participants with irinotecan compared to control (HR 0.74, 95% CI 0.66 to 0.84, high-quality evidence) with low heterogeneity ($I^2 = 0\%$, $P = 0.39$). The study by Boku 2009 alone demonstrated a HR of 0.73 (95% CI 0.64 to 0.83) with median progression-free survival of 4.8 months in the group treated with irinotecan and 4.2 and 2.9 months in the control groups without irinotecan.

Secondary resectability

Information about secondary resectable participants was not provided.

Toxicity

Rates of treatment-related deaths and treatment discontinuation due to toxicity showed substantial heterogeneity between studies ($P = 0.19$, $I^2 = 32\%$) and ($P < 0.00001$, $I^2 = 87\%$), as well as a low event

rate for treatment-related deaths (0.8% versus 1.0%); hence, the significance of pooled results are unclear and will not be discussed in the text of this review (Analysis 5.4; Analysis 5.5). Three studies (Komatsu 2011; Roy 2012; Sugimoto 2014) observed no treatment-related deaths.

Quality of life

Quality of life was assessed in the studies by Bouche 2004, Dank 2008, and Roy 2012. However, in the study by Bouche 2004, which used the EORTC-QLQ C-30 as an assessment tool, the absolute number of participants with follow-up data in the different study arms was very small (between 21 and 29 participants at six months). As compared to treatment with 5-FU/cisplatin, treatment with 5-FU/irinotecan in this study was associated with higher global quality of life and functional scores, as well as lower symptom scores. Dank 2008 compared the time to 5% deterioration of the global health status, as measured by the EORTC QLQ-C30 questionnaire in both treatment arms. In 288 assessable participants (86.5%), the median time to 5% deterioration of the global health status was 4.9 months (95% CI 3.7 to 7.0) in the irinotecan-containing arm and 5.9 months (95% CI 4.8 to 7.7) in the platinum-containing arm. In contrast, the results of the EQ-5D instrument (data from 192 participants): time to definite worsening of Karnofsky performance status, appetite, weight loss, and pain-free survival all favoured the irinotecan-containing arm. Detailed quality-of-life results of this study have been published by Curran 2009. Roy 2012 assessed the clinical benefit in terms of times from baseline to definitive worsening of the Karnofsky performance status (KPS) by at least one category, definitive weight loss by at least 5% and worsening of appetite by at least one grade on a scale of 1 to 5 and added pain-free survival. Median time to definitive deterioration of KPS (4.9 months; 95% CI 1.9 to 11.2 versus 2.6 months; 95% CI not reached) and median time to definitive 5% weight loss (not reached versus 7.6 months) were better without irinotecan. Median time to definitive worsening of appetite was 4.9 months (95% CI not reached) with no difference between groups. Median pain-free survival was not reached in both groups.

(6) Chemotherapy with docetaxel versus non-docetaxel-containing regimens

Overall survival

Eight studies (N = 2001) were summarised in this meta-analysis (Al-Batran 2013; Koizumi 2014; Ochendusko 2015; Ridwelski 2008; Roth 2007; Thuss-Patience 2005; Van Cutsem 2006; Wang 2016). The resulting HR for overall survival was estimated separately for studies with substitutive (i.e. studies in which another chemotherapy was substituted by docetaxel), additive (i.e. studies in which docetaxel was added to other chemotherapies), and other comparisons (Analysis 6.1).

The summary of the three studies (479 participants) with substitutive comparisons (Ridwelski 2008; Roth 2007; Thuss-Patience 2005) slightly favours the non-docetaxel-containing regimens (HR 1.05, 95% CI 0.87 to 1.27, moderate-quality evidence), but does not reach statistical significance. The index of homogeneity according to Thompson ($I^2 = 0\%$) shows a low statistical heterogeneity among the results of these studies (Analysis 6.1). Because necessary data for calculation of the HR in the studies by Sadighi 2006 were missing, this study could not be included in the meta-analysis of this comparison. Differences in pooled median survival between the docetaxel-containing

regimens (9.2 months) and the non-docetaxel-containing regimens (9.4 months) are neither statistically significant nor clinically relevant. All publications describe an adequate allocation concealment and all studies were conducted in Europe.

The meta-analysis of the four studies (N = 1466) (Al-Batran 2013; Koizumi 2014; Van Cutsem 2006; Wang 2016), including a total of 1466 participants, where docetaxel was given in addition to the treatment in the non-docetaxel-containing arm favours the docetaxel-containing regimens (HR 0.80, 95% CI 0.71 to 0.91, moderate-quality evidence) with little heterogeneity between treatment effects of individual studies ($I^2 = 0\%$, $P = 0.82$). This result corresponds to pooled median survival times of 12.3 and 10.6 months, with a small benefit for the docetaxel-containing regimens. Al-Batran 2013 was conducted in Germany, Koizumi 2014 in Japan and Korea, Wang 2016 in China, and the fourth study (Van Cutsem 2006) was an international study that recruited participants in America, Europe, and Asia. Due to the small number of studies in the primary analysis, we did not perform sensitivity analyses.

The single study belonging to other comparisons (Ochendusko 2015) with 56 participants demonstrated a non-statistically significant ($P = 0.43$) advantage of docetaxel-containing regimens compared to control (HR 0.80, 95% CI 0.46 to 1.39, very low-quality evidence). This study permitted second-line therapy with irinotecan monotherapy.

Secondary outcome measures

Tumour response

Data were available from nine eligible studies of 1820 participants (Al-Batran 2013; Dong 2014; Koizumi 2014; Ridwelski 2008; Roth 2007; Sadighi 2006; Thuss-Patience 2005; Van Cutsem 2006; Wang 2016) (Analysis 6.2) (OR 1.37, 95% CI 1.03 to 1.83, moderate-quality evidence).

In studies with substitutive comparisons (Ridwelski 2008; Roth 2007; Sadighi 2006; Thuss-Patience 2005) response rates of 525 participants were 31% in both arms (OR 1.03, 95% CI 0.71 to 1.50, moderate-quality evidence). However, in four studies with additive comparisons and 1235 participants (Al-Batran 2013; Koizumi 2014; Van Cutsem 2006; Wang 2016), pooled response rates of 43% and 30% were observed (OR 1.83, 95% CI 1.45 to 2.32, high-quality evidence). Heterogeneity between results of different studies was very low in both comparisons ($I^2 = 0\%$, $P = 0.98$ and $P = 0.69$, respectively). Among other comparisons, the study by Dong 2014 with 60 participants showed a substantial survival advantage with docetaxel-containing regimens (OR 0.33, 95% CI 0.12 to 0.96, very low-quality evidence).

Time to progression and progression-free survival

Data on time to progression were available for two studies (N = 360) (Ridwelski 2008; Thuss-Patience 2005) (Analysis 6.3) and data for progression-free survival were available for five studies (N = 1498) (Koizumi 2014; Ochendusko 2015; Roth 2007; Van Cutsem 2006; Wang 2016) (Analysis 6.4).

In the case of time to progression, data based on 360 participants from two studies revealed a non-significant difference between docetaxel and non-docetaxel-containing regimens (HR 1.06, 95% CI 0.85 to 1.32, very low-quality evidence) (Analysis 6.3).

In the case of progression-free survival, the pooled hazard ratio was 0.76 (95% CI 0.63 to 0.91, moderate-quality evidence) in favour of docetaxel-containing regimens based on 1498 participants in five studies, but a high level of heterogeneity was observed ($I^2 = 52\%$, $P = 0.08$) (Analysis 6.4). When only additive comparisons were considered (Koizumi 2014; Van Cutsem 2006; Wang 2016) ($N = 1323$), the pooled hazard ratio was 0.70 (95% CI 0.61 to 0.81, high-quality evidence), and a lower level of heterogeneity was observed ($I^2 = 20\%$, $P = 0.29$).

Secondary resectability

This outcome was not reported in any of the studies.

Toxicity

Seven studies ($N = 2113$) reported rates of treatment-related deaths, which were 1.4% in the docetaxel-containing arms versus 1.2% in the non-docetaxel-containing arms (OR 1.10, 95% CI 0.55 to 2.20, moderate-quality evidence, $I^2 = 0\%$, $P = 0.44$) (Analysis 6.5). Treatment discontinuation due to toxicity was numerically less frequent for the participants treated with docetaxel (18.4% versus 21.1%, corresponding to an OR of 0.81 (95% CI 0.53 to 1.25, low-quality evidence), and different results between the studies ($I^2 = 35\%$, $P = 0.19$) (Analysis 6.6). In contrast, discontinuation of treatment due to withdrawal of consent was observed approximately twice as frequently in the participants treated with DCF as compared to CF (22% versus 12%) in the study by Van Cutsem 2006.

Quality of life

Quality of life was assessed by Sadighi 2006, Roth 2007, Van Cutsem 2006, and Al-Batran 2013 with the EORTC-QLQ C30. Furthermore, "clinical benefit", defined as the time to definitive decrease in performance status by \geq one category was analysed in the study by Van Cutsem 2006 and demonstrated a clinical benefit for participants treated with DCF (Ajani 2007).

Sadighi 2006 used the Iranian version of the EORTC-QLQ C30. The report by Ajani 2007a demonstrated a better preservation of Quality of life in the participants treated with DCF. In Roth 2007, global health status improved in participants treated with ECF, but remained stable with both docetaxel regimens. Van Cutsem 2006 measured a longer time to 5% deterioration of the global health status in the docetaxel-containing arm as compared to the non-docetaxel-containing arm (HR 1.44, 95% CI 1.08 to 1.93). Thuss-Patience 2005 measured subjective symptom improvement rates. Al-Batran 2013 (Kripp 2014) administered both EORTC QLQ-C30 and the gastric module STO22, and found that despite the higher toxicity in elderly participants (aged 65 years or older) receiving FLOT, the intensified chemotherapy regimen did not affect quality of life parameters in the elderly.

(7) Chemotherapy with capecitabine versus 5-FU-containing regimens

Overall survival

The results for this comparison are based on 732 randomised participants in five studies (Kang 2009; Li 2016; Ocvirk 2012; Ochendusko 2015; Van Cutsem 2015). The HR for overall survival of 0.94 (95% CI 0.79 to 1.11, moderate-quality evidence) favours the oral regimen, but does not reach statistical significance (Analysis 7.1). The index of heterogeneity for this comparison was small ($I^2 = 12\%$, $P = 0.34$).

The corresponding pooled median survival are 10.8 and 10.9 months, respectively for the capecitabine and 5-FU-containing arms respectively. When the studies which permitted second-line therapy were excluded as part of sensitivity analysis (Li 2016; Ochendusko 2015), the pooled HR for overall survival remained stable at 0.93 (95% CI 0.77 to 1.14).

Secondary outcome measures

Tumour response

Data were available in 636 participants in four studies (Kang 2009; Li 2016; Ocvirk 2012; Van Cutsem 2015). The objective response rate was 38% in both arms, corresponding to an OR of 0.85 (95% CI 0.40 to 1.79, very low-quality evidence) with a non-significant advantage for the capecitabine-containing regimen (Analysis 7.2).

Time to progression

Data from one study of 85 participants (Ocvirk 2012) showed a small benefit for participants treated with capecitabine (HR 0.72, 95% CI 0.47 to 1.12, very low-quality evidence), with improved median time to progression times of 6.8 versus 5.5 months for the participants treated with and without capecitabine.

Kang 2009 provided the largest number of participants for the evaluation of progression-free survival and showed a non-significant advantage for the capecitabine-containing arm (HR 0.80, 95% CI 0.62 to 1.03), corresponding to an improvement in median progression-free survival time of 5.6 versus 5.0 months for the participants treated with and without capecitabine. In total, four studies (Kang 2009; Li 2016; Ochendusko 2015; Van Cutsem 2015) were included in this comparison with a total of 647 participants, demonstrating an overall HR of progression-free survival of 0.98 (95% CI 0.77 to 1.23, very low quality) for participants treated with vs without capecitabine ($I^2 = 23\%$, $P = 0.27$) (Analysis 7.4). The exclusion of studies permitting second-line chemotherapy (Li 2016; Ochendusko 2015) did not alter the findings (HR 0.94, 95% CI 0.58 to 1.53).

Secondary resectability

This outcome was not reported in these studies.

Toxicity

Two studies ($N = 481$) reported deaths due to toxicity, where in the capecitabine-containing arm was 5% in the capecitabine-containing arm and 2% in the 5-FU arm (Kang 2009; Van Cutsem 2015) (Analysis 7.5). The pooled OR for treatment related death is 1.88 (95% CI 0.23 to 15.15, very low-quality evidence), with substantial heterogeneity ($I^2 = 59\%$, $P = 0.12$). Treatment discontinuation due to toxicity was similar in both arms (18%) (Kang 2009) (Analysis 7.6).

Quality of life

Was not reported in these studies.

(8) Chemotherapy with oxaliplatin versus the same regimen including cisplatin

Overall survival

Data were available on 1105 participants in five studies (Al Batran 2008; Hironaka 2016; Kim 2014; Popov 2008; Yamada 2015). The HR for overall survival of 0.81 (95% CI 0.67 to 0.98, low-quality evidence) show a small, advantage for the regimen with oxaliplatin

(Analysis 8.1), with a moderate heterogeneity index ($I^2 = 38\%$, $P = 0.17$). This survival benefit is also reflected in the pooled median overall survival time of 14.0 months versus 11.3 months, respectively.

Secondary outcome measures

Tumour response

Data were available on 1081 participants of the five studies (Al Batran 2008; Hironaka 2016; Kim 2014; Popov 2008; Yamada 2015). The response was 54% in the oxaliplatin-based arms and 47% in the cisplatin-based arms, with a statistically significant advantage for the oxaliplatin-containing regimen (OR 1.38, 95% CI 1.08 to 1.76, moderate-quality evidence), and low heterogeneity between studies ($I^2 = 0\%$, $P = 0.41$) (Analysis 8.2).

Progression-free survival

Progression-free survival, rather than time to progression, was reported in the studies by Al Batran 2008, Hironaka 2016, Kim 2014, and Yamada 2015 ($N = 1034$). The pooled hazard ratio of oxaliplatin-versus cisplatin-based regimens, was 0.88 (95% CI: 0.66 to 1.19) with a high level of heterogeneity between studies ($I^2 = 59\%$, $P = 0.06$) (Analysis 8.3).

Secondary resectability

This outcome was not reported in these studies.

Toxicity

In Popov 2008, two participants (1.4%) experienced a toxic death in the cisplatin-containing arms. These two participants suffered from febrile neutropenia, developed sepsis, and died of septic shock despite antimicrobial therapy. No participant died of toxicity in the oxaliplatin-containing arm (Popov 2008). One treatment-related death occurred in each arm in Kim 2014. Al Batran 2008 observed no treatment-related deaths. Overall, the pooled occurrence of treatment-related deaths (Al Batran 2008; Hironaka 2016; Kim 2014; Popov 2008; Yamada 2015) ($N = 1132$) in both arms were 0.9% and 2.0% in the oxaliplatin- and cisplatin-containing arms, respectively (OR 0.47, 95% CI 0.17 to 1.30, low-quality evidence), with a relatively stable effect across studies ($I^2 = 0\%$, $P = 0.74$) (Analysis 8.4). Data on treatment discontinuation due to toxicity (Al Batran 2008; Hironaka 2016; Yamada 2015) ($N = 970$) were 8% and 10% in the oxaliplatin and cisplatin arms, respectively (OR 0.97, 95% CI 0.44 to 2.13, very low-quality evidence), with substantial between-study heterogeneity ($I^2 = 60\%$, $P = 0.08$) (Analysis 8.5).

Quality of life

This was not reported in these studies.

(9) Taxane-platinum-fluoropyrimidine combinations versus taxane-platinum (without fluoropyrimidine)

Overall survival

The results for this comparison are based on 482 randomised participants in three studies (Ajani 2005; Roth 2007; Van Cutsem 2015). The HR for overall survival of 0.86 (95% CI 0.71 to 1.06, very low-quality evidence) favours the regimen with fluoropyrimidine ($I^2 = 0\%$, $P = 0.47$), but does not reach statistical significance (Analysis 9.1). The corresponding pooled median survival times are 11.7 versus 10.0 months with a small benefit for participants treated

with fluoropyrimidines. Allocation concealment was adequate in the studies included in this comparison.

Secondary outcome measures

Tumour response

Data were available all participants in the above mentioned three studies. The objective response rate was 38% in the 5-FU-containing regimen and 23% in the arm without 5-FU. This corresponds to an OR of 2.08 (95% CI 1.37 to 3.15, low-quality evidence) with an advantage for the 5-FU-containing regimen. Low heterogeneity ($I^2 = 0\%$, $P = 0.89$) between studies was observed.

Time to progression

All three studies reported progression-free survival instead of time to progression. Data from the included studies showed a benefit (HR 0.74 95% CI 0.59 to 0.93, moderate-quality evidence) for the participants treated with 5-FU (Analysis 9.3), and low heterogeneity between studies ($I^2 = 0\%$, $P = 0.83$).

Secondary resectability

This outcome was not reported in these studies.

Toxicity numbers

In three studies ($N = 482$), treatment-related deaths were 6.2% and 2.6% in the 5-FU-containing arms and non-FU-containing arms respectively (OR 1.95, 95% CI 0.73 to 5.17, very low-quality evidence) ($I^2 = 0\%$, $P = 0.88$) (Analysis 9.4). Treatment discontinuation due to toxicity was more frequent for the participants treated with 5-FU (16.7% versus 10.5%), corresponding to an OR of 1.71 (95% CI 0.79 to 3.69, very low-quality evidence) and results between studies were not different ($I^2 = 0\%$, $P = 0.93$) (Analysis 9.5).

Quality of life

This was assessed in only one of these studies (Roth 2007). Treatment burden increased over time in both treatment arms.

(10) S-1 versus 5-FU-containing regimens

Overall survival

This meta-analysis was based on 1793 participants in four randomised studies (Ajani 2010; Boku 2009; Chen 2015; Li 2015). The resulting HR for overall survival of 0.91 (95% CI 0.83 to 1.00, high-quality evidence) demonstrates a borderline statistically significant benefit with low heterogeneity ($I^2 = 0\%$, $P = 0.50$) with overall survival in favour of S-1 (Analysis 10.1). The corresponding pooled median survival times were 9.6 and 9.1 months with a clinically negligible benefit for participants treated with S-1. Allocation concealment was adequate in Ajani 2010 and Boku 2009.

Secondary outcome measures

Tumour response

Data were available in 1753 participants in seven studies (Ajani 2010; Boku 2009; Chen 2015; Dong 2014; Huang 2013; Li 2015; Li 2014). The objective response rate was 32% with S-1 and 26% in the 5-FU arm. This corresponds to an OR of 1.73 (95% CI 1.01 to 2.94, very low-quality evidence) (Analysis 10.2). However, considering the statistically significant heterogeneity ($I^2 = 77\%$, $P = 0.0002$), the results of this comparison have to be evaluated cautiously.

Time to progression

Four studies (Ajani 2010; Boku 2009; Huang 2013; Li 2015) (N = 1942) reported progression-free survival, with a small non-significant benefit for participants treated with S-1 (HR 0.85, 95% CI 0.70 to 1.04, low-quality evidence) (Analysis 10.3) and with substantial heterogeneity ($I^2 = 70\%$, $P = 0.02$).

Three studies (Ajani 2010; Boku 2009; Huang 2013) reported time to treatment failure (TTF) with a HR of 0.84 (95% CI 0.69 to 1.02) in favour of S-1. With the inclusion of Li 2014 and Chen 2015, who reported the time to progression, into the meta-analysis of TTF (Analysis 10.4), (N = 1818), the pooled HR was 0.88 (95% CI 0.76 to 1.01, low-quality evidence), indicating a slight but statistically non-significant benefit of S-1.

Secondary resectability

This outcome was not reported in these studies.

Toxicity

Rates of treatment-related deaths were less frequent in the S-1-containing arm (1.5%) compared to 2.7% in the 5-FU-containing arm (OR 0.56, 95% CI 0.30 to 1.06, moderate-quality evidence) ($I^2 = 0\%$, $P = 0.52$) (Analysis 10.5). Treatment discontinuation due to toxicity was slightly less frequent for the participants treated with S-1 (11.1% versus 12.8%, corresponding to an OR of 0.85 (95% CI 0.63 to 1.13, high-quality evidence) ($I^2 = 11\%$, $P = 0.32$) (Analysis 10.6).

Quality of life

Health-related quality of life was reported by Ajani 2010. There was an advantage of CS compared to CF in terms of the Physical Well-Being (PWB; 51.7% versus 45.1%, $P = 0.044$) component of the FACT-Ga, longer time to worsening of PWB scores (median duration; 4.5 versus 3.0 months, $P = 0.014$), and Chemotherapy Convenience and Chemotherapy Concerns scores.

DISCUSSION

Summary of main results

Only randomised controlled studies were included in this meta-analysis. An exhaustive search for unpublished or ongoing material was performed to minimise publication bias. Sixty studies, with a total of 11,698 participants, have been included in the meta-analysis of overall survival. The meta-analyses of **comparisons 1** (three studies, 184 participants) and **2** (23 studies involving 4447 participants) provide evidence for significant benefits in overall survival for first-line chemotherapy versus best supportive care (BSC), as well as a smaller benefit for combination versus single-agent chemotherapy. Overall, regimens containing irinotecan demonstrated improved results for overall survival in the *substitutive* comparison (i.e. where another chemotherapy was substituted by irinotecan) of irinotecan versus non-irinotecan-containing regimens (six studies with 826 participants, HR 0.87, 95% CI 0.75 to 1.00, high-quality evidence), but not in the additive comparison (i.e. where irinotecan was added to another chemotherapy; HR 0.88, 95% CI 0.76 to 1.03, three studies with 500 participants, low-quality evidence). In contrast, the meta-analysis of all irinotecan-containing versus non-irinotecan-containing regimens demonstrates a small, but significant survival benefit in favour of the people treated with irinotecan. Of note,

both treatment-related deaths and treatment-discontinuation due to toxicity were not increased for people treated with irinotecan.

Furthermore, regimens in which docetaxel was added to a two-drug platinum-fluoropyrimidine combination showed a significant survival benefit (four studies with 1466 participants (HR 0.80, 95% CI 0.71 to 0.91, moderate-quality evidence). In contrast, substituting another chemotherapy (e.g. 5-FU (Ridwelski 2008), epirubicin (Roth 2007), or both epirubicin and 5-FU (Thuss-Patience 2005)) by docetaxel provides no advantage - either in survival, or in secondary outcomes (three studies involving 479 participants, HR 1.05; 0.87 to 1.27, moderate-quality evidence). However, the addition of docetaxel to a two-drug chemotherapy regimen in first-line therapy slightly increases both the risk of treatment-related deaths and treatment discontinuation due to toxicity. The comparison of regimens including capecitabine versus 5-FU showed non-significant advantages in terms of overall survival for the oral 5-FU prodrug capecitabine (732 participants in five studies, 0.94, 95% CI 0.79 to 1.11, moderate-quality evidence).

The comparison of regimens including oxaliplatin versus cisplatin (**comparison 8**), and S-1 versus 5-FU (**comparison 10**), however demonstrated the superior efficacy of oxaliplatin (1105 participants from five studies, HR 0.81, 95% CI 0.67 to 0.98, low-quality evidence) and S-1 (1793 participants in four studies, HR 0.91, 95% CI 0.83 to 1.00, high-quality evidence), respectively, the latter being statistically non significant. However the magnitude of the benefit in individual populations is unclear due to differences in dosing and treatment schedules, and drug metabolism of S-1 between Asian and Caucasian populations. Of note, the landmark REAL-2 study could not be included in this analysis as it included up to 13% of people with oesophageal squamous cell carcinoma in the different study arms.

(1) First-line chemotherapy versus best supportive care

Results for the comparison of first-line chemotherapy versus best supportive care (BSC) convincingly demonstrate a benefit in median survival in favour of chemotherapy (HR 0.37, 95% CI 0.24 to 0.55, three studies, 184 participants, moderate-quality evidence), corresponding to 11 versus 4.3 months weighted average survival. The validity of this result was limited by the small number of participants included in this analysis. In two of four studies addressing this question, either the randomisation (Murad 1993) or the study (Pyrhönen 1995) were terminated early. In another one (Glimelius 1997) the conduct of the study was not possible as planned because the research ethics committee requested the provision of chemotherapy upon request in the BSC group. Further studies addressing this comparison cannot be carried out as BSC cannot be considered to be an appropriate control arm for further studies. Exclusion of the studies by Murad and Pyrhönen, both of which have severe methodological limitations as described above, restricts the studies eligible for this analysis to the study by Scheithauer (Scheithauer 1996). This study, which included 103 participants, was the largest of all studies performed for this comparison and demonstrated a survival benefit of 10.2 versus 5.0 months in the chemotherapy versus BSC group ($P = 0.0001$), which is statistically significant and in line with the results of the other two studies, although not as large. Another study (Glimelius 1997), which was excluded from the analysis (see above) because of cross-over, provided important insights about the quality of life of participants in the chemotherapy and BSC arms. The average quality-adjusted survival was longer in the

group of participants randomised to chemotherapy than in the BSC group (median six versus two months). In addition to the benefits in median survival and quality of life, between 10% and 24% of all participants in the chemotherapy groups in these three studies were alive after two years. In contrast, only one of 81 participants included in the BSC arms of these studies survived longer than 24 months. Two-year survival rates in chemotherapy-treated participants between 5% and 14% were confirmed by other authors (Ohkuwa 2000; Waters 1999), confirming the observation that a limited number of people do have a considerably greater survival benefit from chemotherapy. The reason for the difference in pooled median survival between chemotherapy arms in the studies that compared chemotherapy versus BSC (11.0 months) and the combination therapy arms in studies comparing single-agent versus combination chemotherapy (7.0 months) remains unclear and cannot be explained by differences in prognostic factors. Considering the small number of participants included in the studies comparing chemotherapy versus BSC, as well as the methodological limitations in two of these three studies, an over-estimation of the effect of chemotherapy in these studies is likely.

(2) Single agent versus combination chemotherapy

Regarding **comparison 2**, 16 of 23 relevant individual studies (N = 4447) did not demonstrate a benefit in terms of overall survival for the combination chemotherapy arms. In this context, the results of this meta-analysis demonstrate a statistically significant and consistent benefit for combination versus single-agent therapy in terms of overall survival (HR 0.84, 95% CI 0.79 to 0.89, moderate-quality evidence). Furthermore, response rate and time to progression show advantages for the participants treated with combination chemotherapy. The pooled results of these studies represent a generalised estimate of the effectiveness of the combination chemotherapy regimens used in the last 25 years. Therefore, the benefit of a modern two-drug combination, such as 5-FU/irinotecan or 5-FU/oxaliplatin over a single-agent, usually fluoropyrimidine-based chemotherapy regimen is likely to exceed this global result.

Although any potential survival benefit associated with combination chemotherapy is achieved at the price of increased toxicity, the toxicity of the above mentioned combinations of 5-FU and irinotecan, and 5-FU and oxaliplatin is well managed by oncologists today. Furthermore, given the known correlation between tumour response and quality of life (Sadighi 2006), and considering that the ability of a chemotherapy to maintain a person's health-related quality of life is correlated to its efficacy (Al-Batran 2010), in the absence of contraindications, modern two-drug combination chemotherapy regimens as discussed above should clearly be the preferred option for first-line treatment of people with advanced gastric cancer.

(3) 5-FU/cisplatin/anthracycline combinations versus 5-FU/cisplatin (without anthracyclines)

This comparison was based on 579 participants in four randomised studies and results in a HR of 0.74 (95% CI 0.61 to 0.89, moderate-quality evidence) in favour of the three-drug combination. This comparison is limited by a small number of studies and participants. The results of this comparison are predominantly attributable to the study by Ross 2002, which compared ECF versus MCF. The difference between the data included in this review and the final publication is due to the fact that the original publication

included people with squamous cell cancer of the oesophagus, which do not correspond to the inclusion criteria of this review. Therefore, data from people with gastric adenocarcinoma only as provided by the authors were included in this review.

Whether these results are still relevant today is questionable for two reasons.

1) Since the publication of the REAL-2 study (Cunningham 2008), which demonstrated a significant survival benefit for the combination of EOX (epirubicin, oxaliplatin, and capecitabine) as compared to ECF (epirubicin, cisplatin, and fluorouracil), cisplatin is frequently replaced by oxaliplatin and 5-FU by capecitabine in clinical practice. The relative contribution of epirubicin to the efficacy of the three-drug regimen of capecitabine, oxaliplatin, and epirubicin must be considered as unclear.

2) All studies included in this comparison were conducted at a time when active drugs for second-line chemotherapy were not available. Thus, the validity of these results in 2017, after the publication of 3 randomised studies comparing second-line therapy versus BSC (Ford 2014; Kang 2012; Thuss-Patience 2011), which all three demonstrated as clinically meaningful, statistically significant, and consistent benefit in survival of about 1.5 months, as well as improvements in clinical symptoms Thuss-Patience 2011 and quality of life Ford 2014, needs to be questioned.

In addition, a recently published, french randomised multicentre phase III study Guimbaud 2014, in which a total of 416 participants were included, is of special interest in this context. This study compared the combination of FOLFIRI, followed by ECX (epirubicin, cisplatin, and capecitabine) with the reverse sequence of the same regimens. It has not been included in this meta-analysis as all participants eligible for second-line chemotherapy were systematically crossed over. The results of this study showed similar results for PFS (5.3 versus 5.8 months) and OS (9.5 versus 9.7 months) for both treatment strategies, but a longer time-to-treatment-failure for FOLFIRI. Furthermore, the tolerance of FOLFIRI (overall grade III+IV toxicities and haematological adverse events) was better. For these reasons it must be considered as questionable whether the benefit from adding epirubicin to a two-drug regimen including capecitabine and oxaliplatin outweighs its additional toxicity, especially in sequential treatment strategies.

(4) 5-FU/cisplatin/anthracycline-combinations versus 5-FU/anthracycline combinations (without cisplatin)

Comparison 4 was based on 1147 participants randomised in seven studies and resulted in a HR of 0.82 (95% CI 0.73 to 0.92; low-quality evidence, Analysis 4.1) in favour of the three-drug combination. A sensitivity analysis according to the quality score with inclusion of only those studies in which allocation concealment was adequate (Cocconi 1994; Cocconi 2003; Kikuchi 1990; Webb 1997) does not cause a relevant change of the resulting HR (0.82, 95% CI 0.73 to 0.92, low-quality evidence). Heterogeneity was non-significant in **comparison 3 and 4** ($P = 0.71$ and 0.21 ; I^2 was 0% (95% CI 0% to 26.5%) and 28.5% (95% CI 0% to 69.2%)).

This comparison, which included a greater number of participants and studies compared to **comparison 3**, and is thus much more robust, confirms a statistically significant advantage in overall survival for the addition of cisplatin to the combination of epirubicin and fluorouracil, achieved at the price of increased

toxicity. Again, all studies included in this comparison were published more than 20 years ago, when second-line therapy was unavailable. Therefore, as discussed above, the benefit of anthracyclines in this three-drug combination today is unclear. In view of the evidence discussed above, combinations of 5-FU/cisplatin and an anthracycline are no longer considered as a preferred option for the first-line treatment of advanced gastric cancer today.

(5) Irinotecan versus non-irinotecan-containing regimens

Comparison 5 was based on 2135 participants randomised in 10 studies including six studies in substitutive comparisons, three studies in additive comparison, and two other comparisons. Two treatment arms without irinotecan (5-FU as single-agent and 5-FU/cisplatin) were compared to FOLFIRI in [Bouche 2004](#). Taking all studies into account, the pooled hazard ratio was 0.87 (95% CI 0.80 to 0.95, high-quality evidence) in favour of irinotecan-containing regimens. In subgroup analyses of overall survival, the pooled HR were 0.87 (95% CI 0.75 to 1.00, high-quality evidence) for the substitutive comparison, HR 0.88 (95% CI 0.76 to 1.03, low-quality evidence) for the additive comparison, and HR 0.87 (95% CI 0.76 to 1.00, very low-quality evidence) for the other comparisons, [Analysis 5.1](#).

Objective response rates of 38% versus 30% were observed in substitutive (OR 1.53, 95% CI 0.93 to 2.50, low-quality evidence) and 38% versus 22% in additive comparisons (OR 2.18, 95% CI 1.25 to 3.80, low-quality evidence). The pooled HR for progression-free survival was 0.76 (95% CI 0.69 to 0.84, high-quality evidence); and 0.85 (95% CI 0.72 to 1.00, moderate-quality evidence) and 0.74 (95% CI 0.66 to 0.84, high-quality evidence) for subgroup analysis of substitutive and other comparisons, respectively. Results for rates of treatment-related deaths and treatment discontinuation due to toxicity showed high heterogeneity between studies.

In view of these results, 5-FU/irinotecan-based two-drug combinations should be considered as a true and at least equally effective alternative to platinum-based combinations in first-line therapy. A further advantage of the irinotecan-based combination is the different toxicity profile with no neurotoxicity (as compared to the platinum derivatives) and no significant renal toxicity. In addition, irinotecan-based regimens can easily be administered in the outpatient setting and avoid the hyperhydration necessary for the treatment with cisplatin. Again, the above mentioned study [Guimbaud 2014](#) clearly demonstrates not only the comparable results of treatment with FOLFIRI in first- versus second-line, but as well the feasibility of second-line chemotherapy in 40% to 50% of people and third-line in about 20% of people in Europe.

(6) Docetaxel versus non-docetaxel-containing regimens

Comparison 6 was based on 2001 participants randomised in eight studies including three studies in substitutive comparisons, four studies in additive comparison, and one study in other comparisons. Results from one study ([Sadighi 2006](#)) are not included in this meta-analysis at present because data for the calculation of the HR were not available. Overall, heterogeneity was not significant in the former two comparisons ($I^2 = 0\%$, $P = 0.99$ and 0.82). Of special interest is the fact that studies, in which docetaxel was added to a two-drug regimen of a platinum and a fluoropyrimidine ([Al-Batran 2013](#); [Wang 2016](#); [Van Cutsem 2006](#)) or S-1 as single-agent ([Koizumi 2014](#)) demonstrate a significant benefit not only in terms of survival (HR 0.80; 0.71 to 0.91,

moderate-quality evidence), but also in terms of response rates where the OR was 1.83 (95% CI 1.45 to 2.32, high-quality evidence) for the regimens with docetaxel. In contrast, when docetaxel is substituting another chemotherapy, such as 5-FU, no OS benefit of the docetaxel-containing chemotherapy regimen was observed (HR 1.05, 95% CI 0.87 to 1.27, moderate-quality evidence). Thus, docetaxel-containing two-drug regimens are less efficient than docetaxel-containing three-drug regimens. This observation is confirmed by the recently published, randomised phase II study by [Van Cutsem 2015](#), where overall survival was 14.59 months (95% CI:11.7 to 21.78) for participants treated with docetaxel, oxaliplatin, and 5-FU, as compared to 11.3 (95% CI 8.08 to 14.03) months. However, survival was only a secondary endpoint of this study, and the study was not powered to detect survival differences. Of note, in the study by [Van Cutsem 2006](#) 32% and 41% of participants in both study arms were treated with further chemotherapy lines, while this figure was not given for the study by [Al-Batran 2013](#). The positive effect of the addition of docetaxel to the cisplatin/fluorouracil combination on survival as well as the time to 5% deterioration of global health status in the study by [Van Cutsem 2006](#) was unfortunately achieved at the price of significant toxicity, especially haematological toxicity. For this reason, the clinical value of this regimen is regarded as controversial ([Ilsen 2007](#)). Of note, grade III to IV infection (related to treatment) was more frequent in elderly people (20% versus 9%), and infection was the main cause of treatment-related deaths in both study arms.

Finally, the median age of 55 in the participants included in this study, which was well below the median age of the participants included in other studies (e.g. [Al Batran 2008](#) or [Cunningham 2008](#): 64 and 65 years), needs to be considered when applying these findings to people outside a clinical study. In contrast, the FLOT regimen (5-FU, leucovorin, oxaliplatin, and docetaxel) was developed in an elderly population (median age 69 and 70 years in both treatment groups) ([Al-Batran 2013](#)). The primary endpoint of this randomised phase II study, which compared FLOT with FLO (5-FU, leucovorin and oxaliplatin) was the tolerability and feasibility, defined as per group differences in toxic effects. While the results of this study show - as expected - higher rates of neutropenia, leukopenia, alopecia, and diarrhoea for the participants treated with FLOT, there were similar rates of complicated neutropenia and serious adverse events in the two treatment arms. However, progression-free and overall survival in people over 70 years was similar for treatment with FLOT and FLO. Thus, although FLOT has shown to be feasible in a population over 65 years old, according to a subgroup analysis of younger participants ($n = 68$) versus equal to or older than 70 years ($n = 75$) in this study, only in participants younger than 70 an improved survival was observed for the three-drug combination (median survival 7.1 versus 10.6 months). However, these data are not more than hypothesis generating based on a subgroup analysis from a randomised phase II study. Nevertheless, although the same limitation is valid, another subgroup analysis from this study, which compared the benefit of FLOT to FLO in people with locally advanced versus metastatic disease raised another interesting hypothesis. According to this subgroup analysis, median survival of people with locally advanced disease ($n = 44$) treated with FLOT versus FLO is 24.2 versus 10.3 months, as compared to 7.3 versus 6.0 months in people with metastatic disease ($n = 99$). Thus, people with locally advanced disease might have a greater benefit from FLOT than people with metastatic disease.

(7) Regimens including capecitabine versus intravenous 5-FU-containing regimens

Comparison 7 was based on 732 randomised participants in five studies and resulted in a HR of 0.94 (95% CI 0.79 to 1.11; moderate-quality evidence, [Analysis 7.1](#)). The finding of these studies are in line with the study by [Cunningham 2008](#), which confirms the non-inferiority of capecitabine as compared to 5-FU although it was not included in this comparison because of differences in the participant population (inclusion of people with squamous cell cancer of the oesophagus). For this reason, people with gastric cancer without dysphagia, with adequate renal function and compliance may be treated with capecitabine (or S-1 - see [Analysis 10.1](#)) instead of 5-FU.

(8) Regimens including oxaliplatin versus the same regimen including cisplatin

Overall survival results for **comparison 8** are based on 1105 randomised participants in five studies. The HR for overall survival was 0.81 (95% CI 0.67 to 0.98, low-quality evidence) showing a statistically significant survival advantage in favour of oxaliplatin-containing regimens ([Analysis 8.1](#)). A higher rate of tumour response was also observed in oxaliplatin-containing regimens (54%) compared to cisplatin-containing regimens (47%) (OR 1.38, 95% CI 1.08 to 1.76, moderate-quality evidence). Again, although data from the landmark REAL-2 study ([Cunningham 2008](#)) were not included in this comparison for reasons specified above, they confirm the non-inferiority of oxaliplatin as compared to cisplatin. It is worth noting that three of these included studies were conducted in Asia ([Hironaka 2016](#); [Kim 2014](#); [Yamada 2015](#)), potentially highlighting the applicability of these results to an Asian cohort.

Of special interest in this context is a subgroup analysis of the study by [Al Batran 2008](#), which reports better results for elderly participants treated with oxaliplatin as compared to cisplatin. Therefore, especially when taking into account the higher response rates and lower risk of treatment-related death, oxaliplatin should be preferred to cisplatin in the treatment of gastric cancer.

9) Taxane-platinum-fluoropyrimidine combinations versus taxane-platinum (without fluoropyrimidine)

This comparison was based on 482 randomised participants in three studies and resulted in a HR for overall survival of 0.86 (95% CI 0.71 to 1.06; very low-quality evidence, [Analysis 9.1](#)) in favour of the taxane regimen plus fluoropyrimidine, without reaching statistical significance. The potential drawback was the higher rate of treatment-related deaths (6.2% versus 2.6%, OR 1.95; 95% CI 0.73 to 5.17) and treatment discontinuation due to toxicity (17% versus 11%, OR 1.71, 95% CI 0.79 to 3.69) in the regimens with fluoropyrimidines. Of note; these results are partially attributable to the docetaxel/oxaliplatin/capecitabine-combination, which due to its clearly inferior therapeutic index as compared to the same combination with 5-FU instead of capecitabine ([Van Cutsem 2015](#)) is not recommended. However, objective response rates (38% versus 23%, 2.08 (95% CI 1.37 to 3.15) and progression-free survival (HR 0.74, 95% CI 0.59 to 0.93, moderate-quality evidence) were improved in regimens with fluoropyrimidine. Treatment burden increased over time in both treatment arms. Thus, in conclusion, when a docetaxel-containing three-drug combination chemotherapy regimen is chosen as first-line treatment, oxaliplatin

should be preferred to cisplatin, and 5-FU should be preferred to capecitabine.

(10) S-1 versus 5-FU-containing regimens

This comparison was based on 1793 randomised participants in four studies and resulted in a HR for overall survival of 0.91 (95% CI 0.83 to 1.00; [Analysis 10.1](#)). In addition, a statistically significant advantage in tumour response (OR 1.73, 95% CI 1.01 to 2.94, very low-quality evidence), a numerical benefit in progression-free-survival (HR 0.85, 95% CI 0.70 to 1.04) and time-to-treatment failure (HR 0.88, 95% CI 0.76 to 1.01, both low-quality evidence), less frequent treatment-related deaths (1.5% versus 2.7%, OR 0.56, 95% CI 0.30 to 1.06), and treatment discontinuations due to toxicity (11.1% versus 12.8%, OR 0.85, 95% CI 0.63 to 1.13) were observed in the S-1-containing compared to the 5-FU-containing arms.

Overall completeness and applicability of evidence

In most of these studies, the participants were only in part representative of all people with gastric cancer because they were generally younger than the overall population of people with gastric cancer ([Pye 2001](#)). People with co-morbidities, such as renal or cardiac disease, were excluded. For this reason, these findings are only applicable to people who fulfil the inclusion criteria of these studies and cannot be generalised to all people with gastric cancer. The number of studies representing Asian people with gastric cancer in this 2017 updated review has increased considerably, with now a total of 26 studies conducted at least in part in Asia ([Boku 2009](#); [Chen 2015](#); [Dong 2014](#); [Hironaka 2016](#); [Huang 2013](#); [Kang 2009](#); [Kikuchi 1990](#); [Kim 2001](#); [Kim 2014](#); [Koizumi 2008](#); [Koizumi 2014](#); [Komatsu 2011](#); [KRGGC 1992](#); [Li 2014](#); [Li 2015](#); [Li 2016](#); [Lu 2014](#); [Narahara 2011](#); [Nishikawa 2012](#); [Ohtsu 2003](#); [Shirao 2013](#); [Sugimoto 2014](#); [Wang 2013](#); [Wu 2015](#); [Yamada 2015](#); [Yamamura 1998](#)). The largest number of Asian people was included in comparison 2. Asian people were clearly underrepresented in comparisons 3, 4, and 6. The example of S-1, which is used in different doses in Caucasian (25 mg/m² twice daily) and Asian people (40 mg/m² twice daily) ([Satoh 2014](#)) confirms that chemotherapy regimens need to be tested in Asian and Caucasian populations separately, and that the balance between efficacy and toxicity of a given regimen might be different in different populations due in part to genetic differences ([Syn 2015](#)). Results from clinical studies are thus applicable only to those populations where they have been tested. Except for these limitations, the evidence cited above should be regarded as complete and applicable.

Quality of the evidence

This review included a total of 60 studies and 11,698 participants in the meta-analysis for the primary outcome of overall survival. Seven of the 10 main comparisons for overall survival had low heterogeneity ($I^2 < 20\%$), and even among comparisons with higher levels of inconsistency ($I^2 > 20\%$), the amount of heterogeneity present was not statistically significant ($P > 0.05$). Hence, the pooled results can be considered to be relatively stable. The majority of studies had low risk of bias in terms of random sequence generation, blinding, incomplete efficacy or safety outcome data, and selective reporting. However, the risk of bias due to lack of independent or blinded radiological review and other sources of bias (see [Risk of bias in included studies](#)) are unclear or high in more than 50% of included studies; hence, these can be considered areas for improvement in future studies. The main reasons for

downgrading of evidence in the 'Summary of findings' tables are due to lack of precision in pooled effect sizes, risk of bias (particularly, allocation bias), and statistical heterogeneity.

Potential biases in the review process

For this review, all reasonable effort has been made to reduce and address potential sources of bias, such as inclusion of studies not published in English, searches for unpublished and not fully published studies. Therefore, the likelihood that relevant studies have not been identified is considered as small. One factor with known impact on overall survival after first-line chemotherapy is second-line therapy, which is administered in up to 70% of some recent studies. However, as second-line therapy is now a standard of care, it should not be considered as a source of bias.

Agreements and disagreements with other studies or reviews

Another meta-analysis ([Okines 2008](#)), which summarised the results from two studies that used capecitabine instead of 5-FU, confirmed a significant survival benefit for the people treated with capecitabine, thus lending further support to the use of capecitabine in people with gastric cancer.

We agree with the following key issues in the review by [Garrido 2014](#): DCF is - in terms of efficacy - one of the most promising regimens in younger people with adequate general health. However, it is counterbalanced by significant toxicity, and other three-drug regimens including docetaxel, 5-FU, and oxaliplatin, such as FLOT or TEF ([Van Cutsem 2015](#)) are appropriate alternatives with better tolerability, and that either FOLFIRI or the combination of irinotecan and 5-FU as described by [Dank 2008](#) should also be considered among the most promising regimens on the basis of their significant impact on overall survival, the overall reduced toxicity and time to treatment failure as compared to three-drug regimens ([Guimbaud 2014](#)), as well as the absence of cumulative toxicity. However, we would strongly advise not to use IFL (a combination of bolus 5-FU and irinotecan) in view of the higher rate of treatment-related deaths of this regimen in colorectal cancer ([Hurwitz 2004](#)). The conclusions of the review by [Lordick 2014b](#) "both doublet and triplet drug-regimens can be used.....but careful consideration of the potential toxic complications, impairment of the person's quality of life, and the relative benefit should be undertaken". [Lordick 2014a](#) gives an excellent, more general overview of the current status and challenges in gastric cancer treatment. It addresses not only medical treatment (both chemotherapy and targeted therapies), but also the pathology and surgery. We agree with the main conclusions of the meta-analysis published by the [GASTRIC Group 2013](#) that the addition of experimental chemotherapeutic agents to pre-existing control- or standard regimens have produced a modest improvement in overall survival and progression-free survival, and that none of the regimens emerged as a clear standard. The meta-analysis by [Petrelli 2013](#) compared any two- and three-drug regimens that included CDDP with any regimen containing the same number of agents in which CDDP was replaced by oxaliplatin, CPT-11 or a taxane. We agree with the observation that substitution of cisplatin by modern agents, such as oxaliplatin or irinotecan generally improves outcomes. We also agree with the analysis by ([Chen 2013](#)) that DCF has a better response-rate than non-taxane-containing regimens, but disagree with their statement that

chemotherapy-related toxicity of DCF regimen is acceptable to some extent.

AUTHORS' CONCLUSIONS

Implications for practice

Our review found that people with advanced gastric cancer would benefit from being tested for HER-2 over expression, and in the absence of contraindications receive trastuzumab in combination with cisplatin and 5-FU or capecitabine in case of HER-2 positive disease. For all other people, the use of combination chemotherapy could be considered as standard of care for first-line treatment. The results of this meta-analysis suggest a significant and conclusive survival benefit for chemotherapy versus best supportive care (BSC), as well as a modest survival benefit for combination versus single-agent chemotherapy. As many studies included in this comparison have used combination chemotherapy regimens with suboptimal efficacy, such as 5-FU/epirubicin or 5-FU/ cisplatin, the benefit of combination chemotherapy is likely to be underestimated. In the absence of contraindications, the upfront use of a two-drug combination is efficacious. Among the combination chemotherapy regimens, two-drug combinations including a fluoropyrimidine and oxaliplatin is more efficacious than cisplatin-based combinations in view of their survival benefit and reduced risk of treatment-related deaths. 5-FU/irinotecan-based two-drug combinations are an alternative to platinum-based regimens for first-line treatment, which demonstrated superior results for survival, as well as progression-free survival and tumour response, without any negative impact on toxicity as compared to the non-irinotecan-containing combination chemotherapy regimens. Thus, based on both their efficacy and the balance between efficacy and toxicity, two-drug combinations of a fluoropyrimidine and oxaliplatin or 5-FU and irinotecan are superior treatment regimens for first-line treatment of HER-2 negative gastric cancer. Consideration of the side-effect profile of each regimen is essential in the treatment decision for an individual person.

Three-drug combinations are not widely-used in clinical practice, but might be beneficial for individual people: Two major groups of three-drug-combinations need to be discussed:

1. Docetaxel-based three drug combinations (DCF, FLO-T or TEF).

In which docetaxel is added to a single-agent or two-drug (platinum/5-FU- combination) show significant advantages in terms of overall survival, progression-free-survival, and response rates. However, these advantages are counterbalanced by increased toxicity, especially hematological and neurotoxicity. Of note, while the DCF regimen has been evaluated in a population with a median age of 55 years ([Van Cutsem 2006](#)) with greater toxicity in the elderly people included in this study, it cannot be recommended for elderly people and for this reason FLOT has been shown to be feasible in people over 65 ([Al-Batran 2013](#)). However, in this randomised phase II study conducted in Germany, no benefit from treatment with the three-drug combination of docetaxel, 5-FU, and oxaliplatin (FLOT) as compared to the two-drug regimen FLO (5-FU, leucovorin, and oxaliplatin) was observed in people aged 70 years or older. In the age of personalised medicine, we should not only ask which regimen is better, but which regimen is better for which people. In the above mentioned study by Al-Batran, the benefit from the three-drug combination was limited

to people under 70 years old and a small group of people with locally advanced (versus metastatic) disease. Although data from a subgroup analysis need to be interpreted with caution and the validity of these findings needs to be confirmed in future studies, this is an intriguing hypothesis. Of note, in the perioperative setting an increased pathological response rate was demonstrated for the use of FLOT - as compared to ECF- in a recent publication (Al-Batran 2016), although survival results are still pending.

2. Three-drug regimens which include epirubicin.

Whether the survival benefit for three-drug combinations including cisplatin, 5-FU, and epirubicin - as compared to the same regimen without epirubicin - is still valid when second-line therapy is routinely administered and when cisplatin is replaced by oxaliplatin and 5-FU by capecitabine is questionable.

Furthermore, the magnitude of the observed survival benefits for both groups of three-drug combinations in the palliative setting is not large enough to be clinically meaningful as defined recently by the American Society for Clinical Oncology (ASCO) (Ellis 2014). In contrast to comparisons, in which a survival benefit was observed by adding a third drug to a two-drug regimen at the cost of increased toxicity, the comparison of regimens in which another chemotherapy was replaced by irinotecan was associated with a survival benefit (of borderline statistical significance) *without* increased toxicity.

Nevertheless, individual people in good performance status and with a large tumour burden might have a greater advantage from three-drug regimens (especially docetaxel-containing three-drug combinations) due to their higher response rate.

The benefit of second-line chemotherapy in terms of symptom relief and survival over BSC has been demonstrated in several well-conducted and randomised studies in both European and Asian populations, in people with good performance status. Routine administration of second-line therapy to all people with good performance status might further limit the benefit of a three-drug combination upfront, as compared to the sequential administration of doublets followed by a single-agent, as recently shown in the study by Guimbaud 2014. According to recent phase III studies, the vascular-endothelial-growth-factor-receptor-targeting antibody ramucirumab (Fuchs 2014; Wilke 2014) as single-agent treatment or in combination with paclitaxel chemotherapy might be considered as an alternative to chemotherapy alone as second line.

According to a randomised study among patients with metastatic, non-small cell lung cancer, early palliative care led to significant improvements in both quality of life and mood. As compared to patients receiving standard care, patients receiving early palliative care had less aggressive care at the end of life, but longer survival (Temel 2010). According to their most recent update, ASCO guidelines on "Integration of palliative care into standard oncology care" (Ferrell 2017) recommend that - for newly diagnosed people with advanced cancer - the specialised palliative care team should be involved within eight weeks of diagnosis. Among people with high symptom burdens, outpatient care programs should deliver palliative care services to complement existing program tools. For elderly people, the use of a minimum dataset including the Charlson Comorbidity Index (Charlson 1987), the G8 geriatric assessment screening tool, and the instrumental activities of daily

living should be considered for the assessment of global health status and functional status (Pallis 2011).

Finally, it should be noted that despite all progress in the last decades, advanced gastric cancer remains a disease with a dismal prognosis, and median survival exceeding 12 months only in a small number of randomised studies. Thus, appropriate treatment measure should follow the guidelines of palliative care, defined by WHO (Sepulveda 2002) as "an approach that improves the quality-of-life of people and their families facing the problems associated with life-threatening illness".

Implications for research

Further research is necessary to develop specific treatment strategies for the different subtypes of gastric cancer (TCGA 2014). This concerns not only the molecular subtypes described above, but also clinically defined subgroups of people, for example people with locally advanced or limited metastatic disease, as well as a radiological response to chemotherapy. Considering systemic treatment alone in all people with locally advanced and metastatic gastric cancer is unlikely to achieve the best results for these subgroups with favourable prognostic factors: People with a radiological response in the REAL-2 study (Cunningham 2008) had one- and two-year survival rates of 70.9% (95% CI 61.8 to 78.3) and 27.9% (95% CI 20.0 to 36.2). The value of surgery and/or local ablative therapies in people with good performance status, limited metastatic disease, and respond to chemotherapy is controversial (Kataoka 2017). A randomised clinical study (FLOT-5, NCT02578368) addressing the question if surgery further improves survival and quality of life in people with limited metastatic disease after pretreatment with FLOT is currently ongoing.

The question if strategies to reduce treatment burden, such as maintenance therapy, may be integrated in the continuum of care without compromising efficacy needs to be evaluated in further studies (Digklia 2016).

According to preliminary data from the neoadjuvant setting, people with well differentiated tumours have a greater chance to benefit from chemotherapy than others (Al-Batran 2016). Furthermore, not only targeted therapies, but also chemotherapies might not have the same efficacy in different molecular subtypes of gastric cancer (Tan 2011; Syn 2016)

Further integration of targeted therapies and development of new treatment approaches, such as immunotherapy, for which promising preliminary results have just been presented (Kang 2017; Muro 2016) is highly warranted. Furthermore, treatment strategies combining different immunotherapeutic agents, or chemotherapy and immunotherapy, or immunotherapy and anti-angiogenic therapies need further development. Valid biomarkers, which permit the selection of people with a high chance to respond to the different types of treatments are urgently required.

The role of intraperitoneal chemotherapy, in which cytotoxic agents are infused into the peritoneum, particularly for people with advanced gastric cancer with peritoneal dissemination are being actively investigated (Chan 2017). Randomised studies, including phase III studies, are currently ongoing, and in a future update to this review, it may be worth adding a comparison or subgroup to evaluate the benefits of this mode of chemotherapy administration.

For Her-2 positive breast cancer, great progress has been made with the introduction of agents like pertuzumab and T-DM1 (Swain 2015; Verma 2012), which showed major benefits in recent phase III studies, and many other molecules targeting this pathway are in clinical development. While a benefit for T-DM1 as second-line treatment could not be demonstrated for HER-2 positive gastric cancer (Kang 2016), phase III-results for pertuzumab in gastric cancer are still pending.

For any new treatment, apart from the assessment of outcomes like overall- and progression-free survival, the balance between relief of tumour-associated symptoms and treatment-associated toxicity needs to be evaluated from the person's perspective to determine the palliative value of new therapy regimens in advanced gastric cancer.

Quality of life assessment deserves adequate methodology. Validated measurement instruments are available: a disease-specific module to supplement the EORTC core quality of life questionnaire has been published (Vickery 2001), a FACT-questionnaire for people with stomach cancer has been developed concurrently in North America and Asia (Eremenco 2004), and a clinical meaningful benefit may be defined in advance (Norman 2003). Nevertheless, quality of life assessment remains challenging. Therefore, according to a recent statement from the American Society for Clinical Oncology (Ellis 2014), the interest has shifted to person's self reported specific symptom burden.

An ECOG performance status of 0 to 1 versus 2, and the presence of liver metastases and peritoneal metastases have been identified as poor prognostic factors by multivariate analysis (Chau 2004). A prognostic index was constructed dividing people into a good (no risk factor), moderate (one or two risk factors), and poor (three or more risk factors) risk group. This model has been validated in a second data set from the REAL-2 study. This study did also confirm

the major prognostic impact of obtaining a radiological response: One- and two-year survival rates for responders were 70.9% (95% CI 61.8 to 78.3) and 27.9% (95% CI 20.0 to 36.2). Future studies should explicitly address the value of novel therapeutic strategies in different clinical risk groups.

Importantly, not all questions can be answered in randomised clinical studies. Further research on the risks and benefits of chemotherapy in both the elderly people population, as well as people who do not fulfil the inclusion criteria for clinical studies is required. Therefore, large observational cohort studies or registries of people treated with standard treatment outside clinical studies are important, such as the analysis published by Dixon 2016. Of note, both EORTC and ASCO (Pallis 2011; Wildiers 2013) have published recommendations for designing and reporting clinical studies in Geriatric Oncology in the future, which include obligatory reporting of age-related subgroup analysis, obligatory post-marketing studies in vulnerable and frail older people, and obligatory inclusion of a minimum dataset for senior adults in registration studies and post-marketing studies.

In view of the fact that second-line therapy has become a standard of care in the treatment of this disease, endpoints for clinical studies need reconsideration. Thus, in a disease where the sequential use of different treatments is established, overall survival might not be sufficiently sensitive as an endpoint for clinical studies. However, the correlation of progression-free survival with overall survival in advanced/recurrent gastric cancer was not confirmed in a recent meta-analysis (Paoletti 2013).

ACKNOWLEDGEMENTS

We thank Mrs. Shino Yuo for translations from Japanese and Dr. Yan Pu for translation of Chinese studies, as well as all authors who provided unpublished data.

REFERENCES

References to studies included in this review

Ajani 2005 {published data only}

Ajani JA, Fodor MB, Tjulandin SA, Moiseyenko VM, Chao Y, Cabral FS, et al. Phase II multi-institutional randomized trial of docetaxel plus cisplatin with or without fluorouracil in patients with untreated, advanced gastric, or gastroesophageal adenocarcinoma. *Journal of Clinical Oncology* 2005;**23**(24):5660-7.

Ajani 2010 {published data only}

Ajani JA, Buyse M, Lichinitser M, Gorbunova V, Bodoky G, Douillard JY, et al. Combination of cisplatin/S-1 in the treatment of patients with advanced gastric or gastroesophageal adenocarcinoma: Results of noninferiority and safety analyses compared with cisplatin/5-fluorouracil in the First-Line Advanced Gastric Cancer Study. *European Journal of Cancer* 2013;**49**:3616-24.

* Ajani JA, Rodriguez W, Bodoky G, Moiseyenko V, Lichinitser M, Gorbunova V, et al. Multicenter phase III comparison of cisplatin/S-1 with cisplatin/infusional fluorouracil in advanced gastric or gastroesophageal adenocarcinoma study: the FLAGS trial. *Journal of Clinical Oncology* 2010;**28**(9):1547-53.

Bodoky G, Carrato A, Ravaioli A, Ajani JA. Quality of life in flags trial a randomized, comparative, open label, multicenter, phase 3 of S-1 + cisplatin (CS) compared to 5-fu + cisplatin (CF) in untreated advanced gastric cancer (AGC) patients. *Annals of Oncology*. 2012; Vol. 23 (Supplement 9):ix232-ix233.

Bodoky G, Scheulen ME, Rivera F, Jassem J, Carrato A, Moiseyenko V, et al. Clinical benefit and health-related quality of life assessment in patients treated with cisplatin/S-1 versus cisplatin/5-FU: secondary end point results from the First-Line Advanced Gastric Cancer Study (FLAGS). *Journal of Gastrointestinal Cancer* 2015;**46**(2):109-17.

Al Batran 2008 {published data only}

Al Batran SE, Hartmann JT, Probst S, Schmalenberg H, Hollerbach S, Hofheinz R, et al. Phase III trial in metastatic gastroesophageal adenocarcinoma with fluorouracil, leucovorin plus either oxaliplatin or cisplatin: a study of the Arbeitsgemeinschaft Internistische Onkologie. *Journal of Clinical Oncology* 2008;**26**(9):1435-42.

Al-Batran 2013 {published and unpublished data}

* Al-Batran SE, Pauligk C, Homann N, Hartmann JT, Moehler M, Probst S, et al. The feasibility of triple-drug chemotherapy combination in older adult patients with oesophagogastric cancer: a randomised trial of the Arbeitsgemeinschaft Internistische Onkologie (FLOT65+). *European Journal of Cancer* 2013;**49**(4):835-42.

Kripp M, Al-Batran S E, Rosowski J, Pauligk C, Homann N, Hartmann JT, et al. Quality of life of older adult patients receiving docetaxel-based chemotherapy triplets for esophagogastric adenocarcinoma: a randomized study of the Arbeitsgemeinschaft Internistische Onkologie (AIO). *Gastric Cancer* 2014;**17**:181-7.

Barone 1998 {published data only}

Barone C, Corsi DC, Pozzo C, Cassano A, Fontana T, Noviello MR, et al. Treatment of patients with advanced gastric carcinoma with a 5-fluorouracil-based or a cisplatin-based regimen: two parallel randomized phase II studies. *Cancer* 1998;**82**(8):1460-7. [MEDLINE: 12]

Boku 2009 {unpublished data only}

* Boku N, Yamamoto S, Fukuda H, Shirao K, Doi T, Sawaki A, et al. Fluorouracil versus combination of irinotecan plus cisplatin versus S-1 in metastatic gastric cancer: a randomised phase 3 study. *Lancet Oncology* 2009;**10**(11):1063-9.

Boku N, Yamamoto S, Shirao K, Doi T, Sawaki A, Koizumi W, et al. Randomized phase III study of 5-fluorouracil (5-FU) alone versus combination of irinotecan and cisplatin (CP) versus S-1 alone in advanced gastric cancer (JCOG9912). *Journal of Clinical Oncology, ASCO Annual Meeting Proceedings Part 1* 2007;**25**(18S):LBA4513.

Bouche 2004 {published data only}

Bouche O, Raoul JL, Bonnetain F, Giovannini M, Etienne PL, Lledo G, et al. Randomized multicenter phase II trial of biweekly regime of fluorouracil and leucovorin (LV5FU2), LV5FU2 plus cisplatin, or LV5FU2 plus irinotecan in patients with previously untreated metastatic gastric cancer. Final results of study FFCD 9803. *Journal of Clinical Oncology* 2004;**22**:4319-29.

Cascinu 2011 {published data only}

Cascinu S, Galizia E, Labianca R, Ferrau F, Pucci F, Silva RR, et al. Pegylated liposomal doxorubicin, 5-fluorouracil and cisplatin versus mitomycin-C, 5-fluorouracil and cisplatin for advanced gastric cancer: a randomized phase II trial. *Cancer Chemotherapy and Pharmacology* 2011;**68**(1):37-43.

Chen 2015 {published data only}

Chen JH, Shen WX, Xia JX, Xu RL, Zhu MQ, Xu M. Comparative study between docetaxel, oxaliplatin plus S-1 and DCF regimen as first-line therapy in patients with advanced gastric cancer. *Chinese Journal of Cancer Prevention and Treatment* 2015;**22**(2):134-7.

Cocconi 1994 {published data only}

Cocconi G, Bella M, Zironi S, Algeri R, Di Costanzo F, De Lisi V, et al. Fluorouracil, doxorubicin, and mitomycin combination versus PELF chemotherapy in advanced gastric cancer: a prospective randomized trial of the Italian Oncology Group for Clinical Research. *Journal of Clinical Oncology* 1994;**12**(12):2687-93. [MEDLINE: 23]

Cocconi 2003 {published data only}

Cocconi G, Carlini P, Gamboni A, Gasperoni S, Rodino C, Zironi S, et al. Cisplatin, epirubicin, leucovorin and 5-fluorouracil (PELF) is more active than 5-fluorouracil, doxorubicin and methotrexate (FAMTX) in advanced gastric carcinoma. *Annals of Oncology* 2003;**14**(8):1258-63. [MEDLINE: 2]

Colucci 1995 {published data only}

Colucci G, Giotta F, Maiello E, Cifarelli RA, Leo S, Giuliani F, et al. Efficacy of the association of folinic acid and 5-fluorouracil alone versus folinic acid and 5-fluorouracil plus 4-epidoxorubicin in the treatment of advanced gastric carcinoma. *American Journal of Clinical Oncology* 1995;**18**(6):519-24. [MEDLINE: 24]

Cullinan 1985 {published data only}

Cullinan SA, Moertel CG, Fleming TR, Rubin JR, Krook JE, Everson LK, et al. A comparison of three chemotherapeutic regimens in the treatment of advanced pancreatic and gastric carcinoma: fluorouracil vs fluorouracil and doxorubicin vs fluorouracil, doxorubicin, and mitomycin. *JAMA* 1985;**253**:2061-7.

Cullinan 1994 {published data only}

Cullinan SA, Moertel CG, Wieand HS, O'Connell MJ, Poon MA, Krook JE, et al. Controlled evaluation of three drug combination regimens versus fluorouracil alone for the therapy of advanced gastric cancer. North Central Cancer Treatment Group. *Journal of Clinical Oncology* 1994;**12**(2):412-6. [MEDLINE: 26]

Dank 2008 {published data only}

Curran D, Pozzo C, Zaluski J, Dank M, Barone C, Valvere V, et al. Quality of life of palliative chemotherapy naive patients with advanced adenocarcinoma of the stomach or esophagogastric junction treated with irinotecan combined with 5-fluorouracil and folinic acid: results of a randomised phase III trial. *Quality of Life Research* 2009;**18**:853-61.

* Dank M, Zaluski J, Barone C, Valvere V, Yalcin S, Peschel C, et al. Randomized phase III study comparing irinotecan combined with 5-fluorouracil and folinic acid to cisplatin combined with 5-fluorouracil in chemotherapy naive patients with advanced adenocarcinoma of the stomach or esophagogastric junction. *Annals of Oncology* 2008;**19**(8):1450-7.

De Lisi 1986 {published data only}

De Lisi V, Cocconi G, Tonato M. Randomized comparison of 5-FU alone or combined with carmustine, doxorubicin, and mitomycin (BAFMi) in the treatment of advanced gastric cancer: a phase III trial of the Italian Clinical Research Oncology Group (GOIRC). *Cancer Treatment Reports* 1986;**70**:481-5.

Dong 2014 {published data only}

Dong L, Li J, Lou XP, Miao JH, Lu P, Chang ZW, et al. Comparison of short-term efficacy and safety of TIROX and DCF regimens for advanced gastric cancer. *Journal of International Medical Research* 2014;**42**(3):737-43.

GITSG 1988 {published data only}

Gastrointestinal Tumor Study Group. Triazinate and platinum efficacy in combination with 5-fluorouracil and doxorubicin: results of a three arm randomized trial in metastatic gastric cancer. *Journal of the National Cancer Institute* 1988;**80**:1011-5.

* Koizumi W, Kim YH, Fujii M, Kim HK, Imamura H, Lee KH, et al. Addition of docetaxel to S-1 without platinum prolongs survival of patients with gastric cancer: a randomized study

(START). *Journal of Cancer Research and Clinical Oncology* 2014;**140**:319-28.

Hironaka 2016 {published data only}

Hironaka S, Sugimoto N, Yamaguchi K, Moriwaki T, Komatsu Y, Nishina T, et al. S-1 plus leucovorin versus S-1 plus leucovorin and oxaliplatin versus S-1 plus cisplatin in patients with advanced gastric cancer: a randomised, multicentre, open-label, phase 2 trial. *Lancet Oncology* 2016;**17**(1):99-108. [DOI: 10.1016/S1470-2045(15)00410-6]

Huang 2013 {published data only}

Huang D, Ba Y, Xiong Ji, Xu N, Yan Z, Zhuang Z, et al. A multicentre randomised trial comparing weekly paclitaxel +S-1 with weekly paclitaxel+5-fluorouracil for patients with advanced gastric cancer. *European Journal of Cancer*. England: Department of Gastrointestinal Medical Oncology, Tianjin Medical University Cancer Institute & Hospital, Tianjin, China., 2013; Vol. 49, issue 14:2995-3002.

Kang 2009 {published data only}

Kang YK, Kang WK, Shin DB, Chen J, Xiong J, Wang J, et al. Capecitabine/cisplatin versus 5-fluorouracil/cisplatin as first-line therapy in patients with advanced gastric cancer: a randomised phase III non inferiority trial. *Annals of Oncology* 2009;**20**:666-73. [DOI: 10.1093/annonc/mdn717]

Kikuchi 1990 {published data only}

Kikuchi K, Wakui A, Shimizu H, Kunii Y. Randomized controlled study on chemotherapy with 5-FU, ADM plus CDDP in advanced gastric carcinoma. [Japanese]. *Gan to Kagaku Ryoho [Japanese Journal of Cancer & Chemotherapy]* 1990;**17**(4 Pt 1):655-62. [MEDLINE: 55]

Kim 2001 {published and unpublished data}

Kim TW, Choi SJ, Ahn JH, Bang HS, Chang HM, Kang YK, et al. A prospective randomized phase III trial of 5-fluorouracil and cisplatin (FP) versus epirubicin, cisplatin, and 5-fu (ECF) in the treatment of patients with previously untreated advanced gastric cancer (AGC). *European Journal of Cancer* 2001;**37**(Suppl 6):314. [MEDLINE: 56]

Kim 2014 {published data only}

Kim YS, Sym SJ, Park SH, Park I, Hong J, Ahn HK, et al. A randomized phase II study of weekly docetaxel/cisplatin versus weekly docetaxel/oxaliplatin as firstline therapy for patients with advanced gastric cancer. *Cancer Chemotherapy and Pharmacology* 2014;**73**:163-9.

Koizumi 2008 {published data only}

Koizumi W, Narahara H, Hara T, Takagane A, Akiya T, Takagi M. S-1 plus cisplatin versus S-1 alone for first-line treatment of advanced gastric cancer (SPIRITS trial): a phase III trial. *Lancet Oncology* 2008;**9**(3):215-21.

Koizumi 2014 {published data only}

Fujii M. Chemotherapy for advanced gastric cancer: ongoing phase III study of S-1 alone versus S-1 and docetaxel combination (JACCRO GC03 study). *International Journal of Clinical Oncology* 2008;**13**(3):201-5.

Fujii M, Kim YH, Satoh T, Hosaka H, Kim T, Tsuji A, et al. Randomized phase III study of S-1 alone versus S-1 plus docetaxel (DOC) in the treatment for advanced gastric cancer (AGC): The START trial update. *Journal of Clinical Oncology* 2011;**29**(15 Suppl 1):4016.

Kim YH, Koizumi W, Lee KH, Kishimoto T, Chung HC, et al. Randomized phase III study of S-1 alone versus S-1 plus Docetaxel in the treatment for advanced gastric cancer: The START trial. *Journal of Clinical Oncology*. 2011; Vol. 29:suppl. 4.

* Koizumi W, Kim YH, Fujii M, Kim HK, Imamura H, Lee KH, et al. Addition of docetaxel to S-1 without platinum prolongs survival of patients with advanced gastric cancer: a randomized study (START). *Journal of Cancer Research and Clinical Oncology* 2014;**140**(2):319-28.

Komatsu 2011 {published data only}

Komatsu Y, Takahashi Y, Kimura Y, Oda H, Tajima Y, Tamura S. Randomized phase II trial of first-line treatment with tailored irinotecan and S-1 therapy versus S-1 monotherapy for advanced or recurrent gastric carcinoma (JFMC31-0301). *Anti-cancer drugs* 2011;**22**(6):576-83.

KRGGC 1992 {published data only}

Kyoto Research Group for Chemotherapy of Gastric Cancer. A randomized, comparative study of combination chemotherapies in advanced gastric cancer: 5-fluorouracil and cisplatin (FP) versus 5-fluorouracil, cisplatin, and 4-epirubicin (FPEPIR). *Anticancer Research* 1992;**12**:1983-8.

Levi 1986 {published and unpublished data}

Levi JA, Fox RM, Tattersall MH, Woods RL, Thomson D, Gill G. Analysis of a prospectively randomized comparison of doxorubicin versus 5-fluorouracil, doxorubicin, and BCNU in advanced gastric cancer: implications for future studies. *Journal of Clinical Oncology* 1986;**4**(9):1348-55. [MEDLINE: 70]

Li 2014 {published data only}

Li Y, Zhu D. Comparison between the effect of SOX regimen and FOLFOX4 regimen for advanced gastric cancer. [Chinese] [SOX 方案对比FOLFOX4方案治疗进展期胃癌的临床观察]. *Cancer Research and Clinic* 2014;**26**(1):42-51.

Li 2015 {published data only}

Li YH, Qiu MZ, Xu JM, Sun GP, Lu HS, Liu YP, et al. S-1 plus cisplatin versus fluorouracil plus cisplatin in advanced gastric or gastro-esophageal junction adenocarcinoma patients: a pilot study. *Oncotarget* 2015;**6**(33):35107-15. [DOI: [10.18632/oncotarget.5959](https://doi.org/10.18632/oncotarget.5959)]

Li 2016 {published data only}

Li JY, Huang CZ, Yuan JH, Chen QH. Comparison of efficacy of modified EOX and FOLFIRI regimens in treatment of metastatic gastric cancer. *World Chinese Journal of Digestology* 2016;**24**(12):1866-73.

Loehrer 1994 {published data only}

Loehrer PJ Sr, Harry D, Chlebowski RT. 5-fluorouracil vs. epirubicin vs. 5-fluorouracil plus epirubicin in advanced gastric carcinoma. *Investigational New Drugs* 1994;**12**(1):57-63. [MEDLINE: 72]

Lu 2014 {published data only}

Lu Y, Liu Z, Zhang J. S-1 plus oxaliplatin vs. S-1 as first-line treatment in patients with previously untreated advanced gastric cancer: a randomized phase II study. *Journal of Chemotherapy* 2014;**26**(3):159-64.

Lutz 2007 {published data only}

Lutz MP, Wilke H, Wagener DJ, Vanhoefer U, Jeziorski K, Hegewisch-Becker S, et al. Weekly infusional high-dose fluorouracil (HD-FU), HD-FU plus folinic acid (HD-FU/FA), or HD-FU/FA plus biweekly cisplatin in advanced gastric cancer: randomized phase II trial 40953 of the European Organisation for Research and Treatment of Cancer Gastrointestinal Group and the Arbeitsgemeinschaft Internistische Onkologie. *Journal of Clinical Oncology* 2007;**25**(18):2580-5.

Moehler 2005 {published data only}

Moehler M, Eimermacher A, Siebler J, Höhler T, Wein A, Menges M, et al. Randomized phase II evaluation of irinotecan plus high-dose 5-fluorouracil and leucovorin (ILF) versus 5-fluorouracil, leucovorin and etoposide (ELF) in untreated metastatic gastric cancer. *British Journal of Cancer* 2005;**92**:2122-8.

Moehler 2010 {published data only}

Moehler M, Kanzler S, Geissler M, Raedle J, Ebert MP, et al. A randomized multicenter phase II study comparing capecitabine with irinotecan or cisplatin in metastatic adenocarcinoma of the stomach or esophagogastric junction. *Annals of Oncology* 2010;**21**:71-7. [DOI: [10.1093/annonc/mdp269](https://doi.org/10.1093/annonc/mdp269)]

Murad 1993 {published and unpublished data}

Murad AM, Santiago FF, Petroianu A, Rocha PR, Rodrigues MA, Rausch M. Modified therapy with 5-fluorouracil, doxorubicin, and methotrexate in advanced gastric cancer. *Cancer* 1993;**72**(1):37-41. [MEDLINE: 77]

Narahara 2011 {published and unpublished data}

Chin K, Iishi H, Imamura H, Kobayashi O, Imamoto H, Esaki T, et al. Irinotecan plus S-1 (IRIS) versus S-1 alone as first line treatment for advanced gastric cancer: Preliminary results of a randomized phase III study (GC0301/TOP-002). *Journal of Clinical Oncology, ASCO Annual Meeting Proceedings Part 1* 2007;**25**(18S):4525.

* Narahara H, Iishi H, Imamura H, Tsuburaya A, Chin K, Imamoto H, et al. Randomized phase III study comparing the efficacy and safety of irinotecan plus S-1 with S-1 alone as first-line treatment for advanced gastric cancer (study GC0301/TOP-002). *Gastric Cancer* 2011;**14**(1):72-80.

Nishikawa 2012 {published data only}

Morita S, Baba H, Tsuburaya A, Takiuchi H, Matsui T, Maehara Y, et al. A randomized phase II selection trial in patients with advanced/recurrent gastric cancer: Trial for Advanced Stomach Cancer (TASC). *Japanese Journal of Clinical Oncology* 2007;**37**(6):469-72.

* Nishikawa K, Morita S, Matsui T, Kobayashi M, Takeuchi Y, Takahashi I, et al. A randomized phase-II trial comparing sequential and concurrent paclitaxel with oral or parenteral

fluorinated pyrimidines for advanced or metastatic gastric cancer. *Gastric Cancer* 2012;**15**(4):363-9.

Ochendusko 2015 {published data only}

Ochendusko S, Puskulluoglu M, Konopka K, Fijorek K, Urbanczyk K, Budzynski A, et al. Comparison of efficacy and safety of first-line palliative chemotherapy with EOX and mDCF regimens in patients with locally advanced inoperable or metastatic HER2-negative gastric or gastroesophageal junction adenocarcinoma: a randomized phase 3 trial. *Medical Oncology* 2015;**32**(10):242. [DOI: [10.1007/s12032-015-0687-7](https://doi.org/10.1007/s12032-015-0687-7)]

Ocvirk 2012 {unpublished data only}

Ocvirk J, Rebersek M, Skof E. Randomised prospective phase II study of combination chemotherapy epirubicin, cisplatin, 5-FU (ECF) versus epirubicin, cisplatin, capecitabine (ECX) in patients with advanced or metastatic gastric cancer. *Journal of Clinical Oncology, ASCO Annual Meeting Proceedings Part 1* 2007;**25**(18S):4571.

* Ocvirk J, Rebersek M, Skof E, Hlebanja Z, Boc M. Randomized prospective phase II study to compare the combination chemotherapy regimen epirubicin, cisplatin, and 5-fluorouracil with epirubicin, cisplatin, and capecitabine in patients with advanced or metastatic gastric cancer. *American Journal of Clinical Oncology* 2012;**35**(3):237-41.

Ohtsu 2003 {published and unpublished data}

Ohtsu A, Shimada Y, Shirao K, Bouku N, Hyodo I, Saito H, et al. Randomized phase III trial of fluorouracil alone versus fluorouracil plus cisplatin versus uracil and tegafur plus mitomycin in patients with unresectable, advanced gastric cancer: The Japan Clinical Oncology Group Study (JCOG 9205). *Journal of Clinical Oncology* 2003;**21**(1):54-9. [MEDLINE: 191]

Popov 2002 {published and unpublished data}

Popov I, Svetislav BJ, Jezdic SD. Bi-weekly 24-hour infusion of high dose 5-fluorouracil versus EAP regimen in advanced gastric cancer: a randomised phase II study. *Annals of Oncology*. 2002; Vol. 13 (Suppl. 5):188. [MEDLINE: 150]

* Popov IP, Jelic SB, Krivokapic ZV, Jezdic SD, Pesko PM, Micev MT, et al. Bimonthly 24 h infusion of high-dose 5-fluorouracil vs EAP regimen in patients with advanced gastric cancer. A randomized phase II study. *Medical Oncology* 2008;**25**(1):73-80.

Popov 2008 {published data only}

Popov I, Radosevic-Jelic L, Jezdic S, Milovic M, Borojevic N, Stojanovic S, et al. Biweekly oxaliplatin, fluorouracil and leucovorin versus cisplatin, fluorouracil and leucovorin in patients with advanced gastric cancer. *Journal of the Balkan Union of Oncology* 2008;**13**(4):505-11.

Pyrhonen 1995 {published data only}

Pyrhonen S, Kuitunen T, Nyandoto P, Kouri M. Randomised comparison of fluorouracil, epirubicin and methotrexate (FEMTX) plus supportive care with supportive care alone in patients with non-resectable gastric cancer. *British Journal of Cancer* 1995;**71**(3):587-91. [MEDLINE: 92]

Ridwelski 2008 {unpublished data only}

Ridwelski K, Fahlke J, Schmidt C, Kettner E, Keilholz U, Quitzsch D, et al. Docetaxel-cisplatin (DC) versus 5-fluorouracil-leucovorin-cisplatin (FLC) as first-line treatment for locally advanced or metastatic gastric cancer: Preliminary results of a phase III study. *Journal of Clinical Oncology* 2008;**26** (May 20 Suppl):Abstract 4512.

Ross 2002 {published and unpublished data}

Ross P, Nicolson M, Cunningham D, Valle J, Seymour M, Harper P, et al. Prospective randomized trial comparing mitomycin, cisplatin, and protracted venous-infusion fluorouracil (PVI 5-FU) with epirubicin, cisplatin, and PVI 5-FU in advanced esophagogastric cancer. *Journal of Clinical Oncology* 2002;**20**(8):1996-2004. [MEDLINE: 137]

Roth 1999 {published data only}

Roth A, Kolaric K, Zupanc D, Oresic V, Roth A, Ebling Z. High doses of 5-fluorouracil and epirubicin with or without cisplatin in advanced gastric cancer: a randomized study. *Tumori* 1999;**85**(4):234-8. [MEDLINE: 96]

Roth 2007 {published and unpublished data}

Roth AD, Fazio N, Stupp R, Falk S, Bernhard J, Saletti P, et al. Docetaxel, cisplatin, and fluorouracil; docetaxel and cisplatin; and epirubicin, cisplatin and fluorouracil as systemic treatment for advanced gastric carcinoma: a randomized phase II trial of the Swiss Group for Clinical Cancer Research. *Journal of Clinical Oncology* 2007;**25**(22):3217-23.

Roy 2012 {published data only}

Roy A, Cunningham D, Hawkins R, Sorbye H, Adenis A, Barcelo JR, et al. Docetaxel combined with irinotecan or 5-fluorouracil in patients with advanced oesophago-gastric cancer: a randomised phase II study. *British Journal of Cancer* 2012;**107**(3):435-41.

Sadighi 2006 {published data only}

Sadighi S, Mohagheghi MA, Montazeri A, Sadighi Z. Quality of life in patients with advanced gastric cancer: a randomized trial comparing docetaxel, cisplatin, 5-FU (TCF) with epirubicin, cisplatin, 5-FU (ECF). *BMC Cancer* 2006;**6**:274.

Scheithauer 1996 {published and unpublished data}

Scheithauer W, Kornek G, Hejna M, Depisch D, Raderer M, Huber H. Palliative chemotherapy versus best supportive care in patients with metastatic gastric cancer: a randomized trial. *Annals of Hematology*. 1996; Vol. 73 (Suppl 2):A181. [MEDLINE: 107]

Shirao 2013 {published data only}

Shirao K, Boku N, Yamada Y, Yamaguchi K, Doi T, Goto M, et al. Randomized phase III study of 5-Fluorouracil continuous infusion vs. sequential methotrexate and 5-fluorouracil therapy in far advanced gastric cancer with peritoneal metastasis (JCOG 0106). *Japanese Journal of Clinical Oncology* 2013;**43**(10):972-80.

Sugimoto 2014 {published data only}

Sugimoto N, Fujitani K, Imamura H, Uedo N, Iijima S, Imano M, et al. Randomized phase II trial of S-1 plus irinotecan versus

S-1 plus paclitaxel as first-line treatment for advanced gastric cancer (OGSG0402). *Anticancer Research* 2014;**34**(2):851-7.

Thuss-Patience 2005 {published data only}

Thuss-Patience PC, Kretzschmar A, Repp M, Kingreen D, Henesser D, Micheel S, et al. Docetaxel and continuous infusion fluorouracil versus epirubicin, cisplatin and fluorouracil for advanced gastric adenocarcinoma: a randomized phase II study. *Journal of Clinical Oncology* 2005;**23**(3):494-501.

Van Cutsem 2006 {published data only}

Ajani JA, Moiseyenko VM, Tjulandin S, Majlis A, Constenla M, Boni C, et al. Clinical benefit with docetaxel plus fluorouracil and cisplatin compared with cisplatin and fluorouracil in a phase III trial of advanced gastric or gastroesophageal cancer adenocarcinoma: the V-325 Study Group. *Journal of Clinical Oncology* 2007;**25**:3205-9.

Ajani JA, Moiseyenko VM, Tjulandin S, Majlis A, Constenla M, Boni C, et al. Quality of life with docetaxel plus cisplatin and fluorouracil compared with cisplatin and fluorouracil from a phase III trial for advanced gastric or gastroesophageal adenocarcinoma: the V-325 Study Group. *Journal of Clinical Oncology* 2007;**25**:3210-6.

* Van Cutsem E, Moiseyenko VM, Tjulandin S, Majlis A, Constenla M, Boni C, et al. Phase III study of docetaxel and cisplatin plus fluorouracil compared with cisplatin and fluorouracil as first-line therapy for advanced gastric cancer: a report of the V325 Study Group. *Journal of Clinical Oncology* 2006;**24**(31):4991-7.

Van Cutsem 2015 {published data only}

Van Cutsem E, Boni C, Tabernero J, Massuti B, Middleton G, Dane F, et al. Docetaxel plus oxaliplatin with or without fluorouracil or capecitabine in metastatic or locally recurrent gastric cancer: a randomized phase II study. *Annals of Oncology* 2015;**26**(1):149-56.

Wang 2013 {published data only}

Wang X, Wang ML, Zhou LY, Lu XY, Yang JF, Yu HG. Randomized phase II study comparing paclitaxel with S-1 vs. S-1 as first-line treatment in patients with advanced gastric cancer. *Clinical and Translational Oncology*. Springer Milan (Via Podgora 4, Milan I-20122, Italy), 2013; Vol. 15, issue 10:836-42.

Wang 2016 {published data only}

Wang J, Xu R, Li J, Bai Y, Liu T, Jiao S, et al. Randomized multicenter phase III study of a modified docetaxel and cisplatin plus fluorouracil regimen compared with cisplatin and fluorouracil as first-line therapy for advanced or locally recurrent gastric cancer. *Gastric Cancer* 2016;**19**(1):234-44.

Webb 1997 {published and unpublished data}

Webb A, Cunningham D, Scarffe JH, Harper P, Norman A, Joffe JK, et al. Randomized trial comparing epirubicin, cisplatin, and fluorouracil versus fluorouracil, doxorubicin, and methotrexate in advanced esophagogastric cancer. *Journal of Clinical Oncology* 1997;**15**(1):261-7. [MEDLINE: 148]

Wu 2015 {published data only}

Wu D, Li X, Tong J, Sun L, Zheng H, Gao C, et al. S-1 combined with cisplatin versus cisplatin alone for the treatment of advanced gastric cancer: a pilot randomized-controlled trial. *Anticancer Drugs* 2015;**26**(7):774-8. [DOI: [10.1097/CAD.0000000000000242](https://doi.org/10.1097/CAD.0000000000000242)]

Yamada 2015 {published data only}

Yamada Y, Higuchi K, Nishikawa K, Gotoh M, Fuse N, Sugimoto N, et al. Phase III study comparing oxaliplatin plus S-1 with cisplatin plus S-1 in chemotherapy-naïve patients with advanced gastric cancer. *Annals of Oncology* 2015;**26**:141-8.

Yamamura 1998 {published and unpublished data}

Yamamura Y, Miyazaki I, Ogawa M, Yonemura Y, Tanemura H, Kito T, et al. A randomized controlled trial with methotrexate (MTX), 5-fluorouracil (5-FU) and pirarubicin (THP) vs 5-FU alone in advanced or recurrent gastric carcinoma. Tokai Hokuriku THP Study Group. [Japanese]. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1998;**25**(10):1543-8. [MEDLINE: 132]

References to studies excluded from this review

Ahn 2002 {published and unpublished data}

Ahn JH, Kang YK, Kim TW, Bahng H, Chang HM, Kang WC, et al. Nephrotoxicity of heptaplatin: a randomized comparison with cisplatin in advanced gastric cancer. *Cancer Chemotherapy & Pharmacology* 2002;**50**(2):104-10. [MEDLINE: 2]

Ajani 2002 {published data only}

Ajani JA. Docetaxel in combination for advanced gastric cancer. *Gastric Cancer* 2002;**5**(Suppl 1):31-4. [MEDLINE: 5]

Ajani 2006 {published data only}

Ajani JA, Lee F-C, Singh DA, Haller DG, Lenz H-J, Benson AB III, et al. Multicenter phase II trial of S-1 plus cisplatin in patients with untreated advanced gastric of gastroesophageal junction adenocarcinoma. *Journal of Clinical Oncology* 2006;**24**(4):663-7.

Akagi 2010 {published data only}

Akagi J, Baba H. PSK may suppress CD57(+) T cells to improve survival of advanced gastric cancer patients. *International Journal of Clinical Oncology* 2010;**15**(2):145-52.

Akazawa 1985 {published data only}

Akazawa S, Nakajima T, Kitagawa H, Nakagawa T, Kanda Y, Futatsuki K, et al. Therapeutic effect of sequential doses of methotrexate (MTX) and 5-fluorouracil (5-FU) in advanced gastric cancer: comparison of intermediate-dose MTX with high-dose MTX. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1985;**12**(1):91-8. [MEDLINE: 140]

Andrić 2012 {published data only}

Andrić Z, Randjelović T, Kovčin V, Gutović J, Crevar S, Murtezani Z, et al. Evaluation of the efficacy and toxicity of protocol cisplatin, 5-fluorouracil, leucovorin compared to protocol fluorouracil, doxorubicin and mitomycin C in locally advanced and metastatic gastric cancer. *Srpski Arhiv za Celokupno Lekarstvo* 2012;**140**(5-6):305-12.

Anonymous 1979 {published data only}

Anonymous. Phase II-III chemotherapy studies in advanced gastric cancer. The Gastrointestinal Tumor Study Group. *Cancer Treatment Reports* 1979;**63**(11-12):1871-6. [MEDLINE: 3]

Anonymous 1982 {published data only}

* Anonymous. A comparative clinical assessment of combination chemotherapy in the management of advanced gastric carcinoma: The Gastrointestinal Tumour Study Group. *Cancer* 1982;**49**(7):1362-6.

O'Connell MJ, O'Fallon Y, Lavin PT, Moertel CG, Bruckner HW, Douglass HO Jr, et al. A comparative assessment of combination chemotherapy in advanced gastric cancer. Proceedings/Annual Meeting of the American Society of Clinical Oncology. 1980:Abstract 403. [MEDLINE: 85]

Anonymous 1983 {published data only}

Anonymous. Chemotherapy for advanced stomach cancer - a controlled study of AF and MF. [Japanese]. *Gan to Kagaku Ryoho [Japanese Journal of Cancer & Chemotherapy]* 1983;**10**(10):2171-8. [MEDLINE: 5]

Anonymous 1984 {published data only}

* Anonymous. Randomized study of combination chemotherapy in unresectable gastric cancer. The Gastrointestinal Tumor Study Group. *Cancer* 1984;**53**(1):13-7.

O'Connell DM, Stablein DM for the Gastrointestinal Tumor Study Group. A prospective clinical trial of 5-fluorouracil/adriamycin based chemotherapy in unresectable gastric cancer. Proceedings/Annual Meeting of the American Society of Clinical Oncology. 1982; Vol. 2:91. [MEDLINE: 86]

Aoyama 1981 {published data only}

Aoyama M, Hirose H, Adachi N. Comparison of combination therapy of 5-fluorouracil, mitomycin C, and adriamycin (FAM) and mitomycin C, 5-fluorouracil and cytosine arabinoside (MFC) for advanced gastric cancer. [Japanese]. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1981;**8**(5):757-62. [MEDLINE: 9]

Bajetta 1998 {published data only}

Bajetta E, Di Bartolomeo M, Carnaghi C, Buzzoni R, Mariani L, Gebbia V, et al. FEP regimen (epidoxorubicin, etoposide and cisplatin) in advanced gastric cancer, with or without low-dose GM-CSF: an Italian Trial in Medical Oncology (ITMO) study. *British Journal of Cancer* 1998;**77**(7):1149-54. [MEDLINE: 10]

Baker 1976 {published data only}

Baker LH, Talley RW, Matter R, Lehane DE, Rutter BW, Jones SE, et al. Phase III comparison of the treatment of advanced gastrointestinal cancer with bolus weekly 5-FU vs. methyl-CCNU plus bolus weekly 5-FU. *Cancer* 1976;**38**:1-7. [MEDLINE: 163]

Balana 1990 {published data only}

Balana C, Camps C, Diaz Rubio E, Jimeno J, Dorta J, Massuti B, et al. Treatment of advanced gastric cancer (GC) with a fluorouracil (F) or a cisplatin (P) based chemotherapy. A study of the Spanish cooperative group for gastrointestinal tumor therapy (TTD). *Annals of Oncology*. 1990; Vol. 1 (Suppl):43. [MEDLINE: 11]

Berenberg 1989 {published data only}

Berenberg JL, Goodman PJ, Oishi N, Fleming T, Natale RB, Hutchins LH, et al. [5-Fluorouracil (5-FU) and folinic acid (FA): for the treatment of metastatic gastric cancer]. Proceedings/Annual Meeting of the American Society of Clinical Oncology. 1989; Vol. 8:101.

Berenberg 1995 {published data only}

Berenberg JL, Tangen C, Macdonald JS, Hutchins LF, Natale RB, Oishi N, et al. Phase II study of 5-fluorouracil and folinic acid in the treatment of patients with advanced gastric cancer. A Southwest Oncology Group Study. *Cancer* 1995;**76**(5):715-9. [MEDLINE: 136]

Beretta 1983 {published data only}

Beretta G, Fracchini P, Ravaioli A, Amadori D, Luporini G. FAM/FAMB Polychemotherapy for advanced carcinoma of the stomach (ACS): a randomized study. Proceedings/Annual Meeting of the American Society of Clinical Oncology. 1983; Vol. 2:131. [MEDLINE: 159]

Beretta 1989 {published data only}

Beretta G, Arnoldi E, Beretta GD, Tedeschi L, Dallavalle G, Bollina R, et al. A randomized study of fluorouracil versus FAM polychemotherapy in gastric carcinoma. Proceedings of the EORTC Symposium on Advances in Gastrointestinal Tract Cancer Research and Treatment. Strasbourg, 1989:P.48. [MEDLINE: 162]

Berglund 2006 {published data only}

Berglund A, Byström P, Pedersen D, Nygren P, Frödin JE, Bergman A, et al. GI-TAC: A randomised phase II study of sequential docetaxel and irinotecan with 5-fluorouracil, folinic acid in patients with metastatic upper abdominal (pancreatic, gastric or biliary) cancer. *Annals of Oncology* 2006;**17**(Suppl 9: Abstract 1126P):ix308-ix326.

Bi 2011 {published data only}

Bi F, Li Q, Zhou C, Yu J, Cai X, Qiu M, et al. Preliminary results of a randomized phase II study: Treatment of Chinese patients with advanced gastric cancer with FOLFIRI followed by FOLFOX7 or the reverse sequence. *Journal of Clinical Oncology*. 2011; Vol. 29 (Supp 1):15.

Bjerkset 1986 {published data only}

Bjerkset T, Fjosne HE. Comparison of oral ftorafur and intravenous 5-fluorouracil in patients with advanced cancer of the stomach, colon or rectum. *Oncology* 1986;**43**(4):212-5. [MEDLINE: 13]

Bruckner 1986 {published data only}

Bruckner HW, Stablein DM for the Gastrointestinal Tumor Study Group. A randomized study of 5-fluorouracil (F) and doxorubicin (A) with semustine (Me), cis-platinum (P), or triazine (T) for treatment of advanced gastric cancer. Proceedings/Annual Meeting of the American Society of Clinical Oncology. 1986; Vol. 5:90. [MEDLINE: 164]

Brugarolas 1975 {published and unpublished data}

Brugarolas A, Garcia MM, Lacave AJ. Chemotherapy in advanced gastric cancer. A controlled clinical study. Proceedings /Annual

Meeting of the American Association for Cancer Research. 1975; Vol. 16(66):169. [MEDLINE: 15]

Bugat 2003 {published data only}

Bugat R. Irinotecan in the treatment of gastric cancer. *Annals of Oncology* 2003;**14**(Suppl 2):37-40. [MEDLINE: 13]

Buroker 1979 {published data only}

Buroker T, Kim PN, Gropp C, McCracken J, O'Bryan R, Panettiere F, et al. 5 FU infusion with mitomycin-C vs. 5 FU infusion with methyl-CCNU in the treatment of advanced upper gastrointestinal cancer: a Southwest Oncology Group Study. *Cancer* 1979;**44**(4):1215-21. [MEDLINE: 16]

Cai 2011 {published data only}

Cai X, Xue P, Song WF, Hu J, Gu HL, Yang HY, et al. The role of pharmacokinetic monitoring of fluorouracil in improvement of efficacy and reduction of adverse reactions for patients with advanced gastric cancer. *Tumor* 2011;**31**(10):930-6.

Cascinu 1994 {published data only}

Cascinu S, Cordella L, Catalano G. Neuroprotective effect of reduced glutathione (GSH) on cisplatin based chemotherapy in advanced gastric cancer: a double blind randomized trial. Proceedings/Annual Meeting of the American Society of Clinical Oncology. 1994; Vol. 13:431. [MEDLINE: 17]

Cascinu 1995 {published data only}

Cascinu S, Cordella L, Del Ferro E, Fronzoni M, Catalano G. Neuroprotective effect of reduced glutathione on cisplatin-based chemotherapy in advanced gastric cancer: a randomized double-blind placebo-controlled trial. *Journal of Clinical Oncology* 1995;**13**(1):26-32. [MEDLINE: 18]

Cascinu 1996 {published data only}

Cascinu S, Del Ferro E, Catalano G. Different doses of granulocyte colony stimulating factor to support a weekly chemotherapeutic regimen in advanced gastric cancer: a randomized study. *Anti-Cancer Drugs* 1996;**7**(1):43-7. [MEDLINE: 19]

Chau 2013 {published data only}

Chau I, Passalacqua R, Zalberg JR, Fuchs CS, Liepa AM, Hsu Y, et al. Tolerability and quality-of-life (QoL) results from the phase 3 REGARD study: Ramucirumab versus placebo in patients with previously treated gastric or gastroesophageal junction (GEJ) adenocarcinoma. *European Journal of Cancer*. Elsevier Ltd, 2013; Vol. 49:S615.

Chen 2011 {published data only}

Chen D, Jia R, Gao S, Feng X, Pan L, Song N, et al. Effect of low-dose gimeracil and oteracil potassium combined with oxaliplatin on aged patients with advanced gastric cardiac adenocarcinoma. *Chinese Journal of Clinical Oncology* 2011;**38**(7):396-9.

Chlebowski 1979 {published data only}

Chlebowski RT, Paroly WS, Pugh RP, Weiner JM, Bateman JR. Treatment of advanced gastric carcinoma with 5-fluorouracil: a randomized comparison of two routes of delivery. *Cancer Treatment Reports* 1979;**63**(11-12):1979-81. [MEDLINE: 138]

Chlebowski 1985 {published data only}

Chlebowski RT, Weiner JM, Silverberg I, Glass A, Bateman JR. Cyclophosphamide plus 5-FU versus 5-FU alone in advanced gastric carcinoma. *Oncology* 1985;**42**(3):141-3. [MEDLINE: 20]

Chu 2006 {published data only}

Chu JH, Zhang Y, Liu DF, Ji HM. Weekly docetaxel combined with cisplatin of fluorouracil for advanced gastric carcinoma [Chinese]. *Chinese Journal of Clinical Oncology* 2006;**11**:541-2.

Chung 2011 {published data only}

Chung KY, Saito K, Zergebel C, Hollywood E, Segal M, Saltz LB. Phase I study of two schedules of oral S-1 in combination with fixed doses of oxaliplatin and bevacizumab in patients with advanced solid tumors. *Oncology* 2011;**8**(2):65-72.

Coates 1984 {published data only}

Coates AS, Tattersall MH, Swanson C, Hedley D, Fox RM, Raghavan D. Combination therapy with methotrexate and 5-fluorouracil: a prospective randomized clinical trial of order of administration. *Journal of Clinical Oncology* 1984;**2**(7):756-61. [MEDLINE: 21]

Cocconi 1982 {published data only}

Cocconi G, DeLisi V, Di Blasio B. Randomized comparison of 5-FU alone or combined with mitomycin and cytarabine (MFC) in the treatment of advanced gastric cancer. *Cancer Treatment Reports* 1982;**66**(6):1263-6. [MEDLINE: 22]

Cocconi 1992 {published data only}

Cocconi G, Bella M, Zironi S, Algeri R, Bartolucci R, De Lisi V, et al. A prospective randomized trial comparing FAM combination with PELF combination in advanced gastric carcinoma. Proceedings/Annual Meeting of the American Society of Clinical Oncology. 1992; Vol. 11:163. [MEDLINE: 166]

Colucci 1991 {published data only}

Colucci G, Maiello E, Valori V, Pezzella G, Giuliani F. A randomized study with 5-FU + folinic acid (FA) alone or with epirubicin (E) in advanced gastric carcinoma. *European Journal of Cancer*. 1991; Vol. 27 Suppl. 2:S81. [MEDLINE: 168]

Constenla 2002 {published data only}

Constenla M, Garcia-Arroyo R, Lorenzo I, Carrete N, Campos B, Palacios P. Docetaxel, 5-fluorouracil, and leucovorin as treatment for advanced gastric cancer: results of a phase II study. *Gastric Cancer* 2002;**5**(3):142-7. [MEDLINE: 7]

Coombes 1994 {published data only}

Coombes RC, Chilvers CE, Amadori D, Medi F, Fountzilias G, Rauschecker H, et al. Randomised trial of epirubicin vs fluorouracil in advanced gastric cancer. An International Collaborative Cancer Group (ICCG) study. *Annals of Oncology* 1994;**5**:33-6. [MEDLINE: 189]

Cullinan 1993 {published data only}

Cullinan S, Moertel CG, Wieand H, Poon M. A randomized comparison of fluorouracil + adriamycin + cisplatin (FAP); fluorouracil + adriamycin + seumustine (FAME), FAME alternating with triazinate (T), and fluorouracil alone in advanced gastric carcinoma. A North Central Cancer Treatment

Group Study. Proceedings/Annual Meeting of the American Society of Clinical Oncology. 1993; Vol. 12:200. [MEDLINE: 169]

Cunningham 2008 {published data only}

Cunningham D, Starling N, Rao S, Iveson T, Nicolson M, Coxon F, et al. Capecitabine and oxaliplatin for advanced esophagogastric cancer. *New England Journal of Medicine* 2008;**358**:36-46.

De Lisi 1985 {published data only}

De Lisi V, Tonato M, Leonardi F, Soldani M, Di Constanzo F, Marinelli G, et al. Randomized comparison of 5-fluorouracil (5-FU) versus a combination of BCNU (B), adriamycin (A), 5-FU and mitomycin C (Mi) (BAFMI) in the treatment of advanced gastric carcinoma. Proceedings/Annual Meeting of the American Association for Cancer Research. 1985; Vol. 26:696. [MEDLINE: 27]

De Lisi 1988 {published data only}

De Lisi V, Cocconi G, Tonato M, Di Costanzo F, Leonardi F, Soldani M. 5-Fluorouracil versus a combination of BCNU, adriamycin, 5-FU and mitomycin C in advanced gastric cancer: a prospective randomized study of the Italian Clinical Research Oncology Group. *Recent Results in Cancer Research* 1988;**110**:212-8. [MEDLINE: 29]

Diaz-Rubio 1991 {published data only}

Diaz-Rubio E, Jimeno J, Cervantes A, Cruz J, Camps C, Massutic B, et al. Treatment of stage IV gastric cancer (GC) with a fluorouracil (F) or a cisplatin (P) based combination chemotherapy. Proceedings/Annual Meeting of the American Society of Clinical Oncology. 1991; Vol. 10:158. [MEDLINE: 171]

Douglass 1983 {published data only}

Douglass HO Jr, Lavin PT, Goudsmit A, Klaassen DJ, Paul AR. Phase II-III evaluation of combinations of methyl-CCNU, mitomycin C, adriamycin, and 5-fluorouracil in advanced measurable gastric cancer. Proceedings/Annual Meeting of the American Society of Clinical Oncology. 1983; Vol. 2:121. [MEDLINE: 161]

Douglass 1984 {published data only}

Douglass HO Jr, Lavin PT, Goudsmit A, Klaassen DJ, Paul AR. An Eastern Cooperative Oncology Group evaluation of combinations of methyl-CCNU, mitomycin C, adriamycin, and 5-fluorouracil in advanced measurable gastric cancer (EST 2277). *Journal of Clinical Oncology* 1984;**2**(12):1372-81. [MEDLINE: 32]

Duffour 2006 {published data only}

Duffour J, Bouche O, Rougier P, Milan C, Bedenne L, Seitz JF, et al. Safety of cisplatin combined with continuous 5-FU versus bolus 5-FU and leucovorin, in metastatic gastrointestinal cancer (FFCD 9404 randomised trial). *Anticancer Research* 2006;**26**(5B):3877-83.

Figoli 1991 {published data only}

Figoli F, Galligioni E, Crivellari D, Frustaci S, Talamini R, Sorio R, et al. Evaluation of two consecutive regimens in advanced gastric cancer. *Cancer Investigation* 1991;**9**(3):257-62. [MEDLINE: 34]

Ford 2014 {published data only}

Ford HER, Marshall A, Bridgewater JA, Janowitz T, Coxon FY, Wadsley J, et al. Docetaxel versus active symptom control for refractory oesophagogastric adenocarcinoma (COUGAR-02): an open-label, phase 3 randomised controlled trial. *Lancet Oncology* 2014;**15**(1):78-86.

Fuchs 2014 {published data only}

Fuchs CS, Tomasek J, Yong CJ, Dumitru F, Passalacqua R, Goswami C, et al. REGARD Trial Investigators. Ramucirumab monotherapy for previously treated advanced gastric or gastro-oesophageal junction adenocarcinoma (REGARD): an international, randomised, multicentre, placebo-controlled, phase 3 trial [DW11]. *Lancet*. England: Department of Medical Oncology, Dana-Farber Cancer Institute, Harvard Medical School, Boston, MA, USA. Electronic address: cfuchs@partners.org., 2014; Vol. 383, issue 9911:31-9.

Fujii 1983 {published data only}

Fujii M, Murakami N. Single chemotherapy with 5-fluorouracil or fluorouracil and combination chemotherapy with mitomycin C and 5-fluorouracil for inoperable elderly gastric cancers. [Japanese]. *Nihon Gan Chiryō Gakkai Shi* 1983;**18**(1):1-4. [MEDLINE: 36]

Furue 1985 {published data only}

Furue H, Uchino H, Orita K, Kimura T, Goto Y, Kondo T, et al. Clinical evaluation of schizophyllan (SPG) in advanced gastric cancer (the second report) - a randomized controlled study. [Japanese]. *Gan to Kagaku Ryōho. Cancer & Chemotherapy* 1985;**12**(6):1272-7. [MEDLINE: 37]

Furukawa 1995 {published data only}

Furukawa H, Iwanaga T, Nakajima T, Okabayashi K, Nakazato H, Hiratsuka M, et al. Randomized study with mitomycin C+ 5-fluorouracil+cytosine arabinoside (MFC)+5-fluorouracil, MFC + tegafur and uracil (UFT), and MF+UFT in advanced gastric cancer: Interinstitutional differences in a Multicenter Study in Japan. *Journal of Surgical Oncology* 1995;**60**(1):59-64. [MEDLINE: 38]

Gao 2010 {published data only}

Gao H, Ding X, Wei D, Xu T, Cheng P. Docetaxel versus epirubicin in combined with cisplatin, leucovorin and fluorouracil for advanced gastric carcinoma as first line therapy: a randomized clinical trial. *Chinese Journal of Clinical Oncology* 2010;**15**:529-33.

Gioffre 1984 {published data only}

Gioffre FM, Caputo G, Lucania L. 5-FU + BCNU versus FAM for advanced gastric cancer. [Italian]. *Folia Oncologica* 1984;**7**(3):198-203. [MEDLINE: 40]

Glimelius 1994 {published data only}

Glimelius B, Hoffman K, Haglund U. Initial or delayed chemotherapy with best supportive care in advanced gastric cancer. *Annals of Oncology* 1994;**5**:189-90. [MEDLINE: 187]

Glimelius 1995 {published data only}

Glimelius B, Hoffman K, Graf W, Haglund U, Nyren O, Pahlman L, et al. Cost-effectiveness of palliative chemotherapy in advanced

gastrointestinal cancer. *Annals of Oncology* 1995;**6**(3):267-74. [MEDLINE: 41]

Glimelius 1997 {published data only}

Glimelius B, Ekstrom K, Hoffman K, Graf W, Sjoden PO, Haglund U, et al. Randomized comparison between chemotherapy plus best supportive care with best supportive care in advanced gastric cancer. *Annals of Oncology* 1997;**8**(2):1-6. [MEDLINE: 42]

Goseki 1995 {published data only}

Goseki N, Maruyama M, Nagai K, Kando F, Endo M, Shimoju K, et al. Clinical evaluation of anticancer effect of methionine-depleting total parenteral nutrition with 5-fluorouracil and/or mitomycin C. [Japanese]. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1995;**22**(8):1028-35. [MEDLINE: 43]

Grau 1988 {published data only}

Grau JJ, Estapé J, Alcobendas F, Gracia J, Antón A, Cruz JJ, et al. Mitomycin C-florafur versus mitomycin C alone as complementary chemotherapy in locally advanced gastric cancer. *Neoplasia* 1988;**5**:179-81. [MEDLINE: 45]

Grieco 1984 {published data only}

Grieco A, Lasorella A, Astone A, Vagliviello L, Bartoloni C, Barone C. Chemotherapy of gastrointestinal cancer in the advanced stage with a combination of 5-fluorouracil, adriamycin, mitomycin C (FAM). A non-controlled study [Italian]. *Clinica Terapeutica* 1984;**110**(4):353-8. [MEDLINE: 46]

Gubanski 2010 {published data only}

Gubanski M, Johnsson A, Fernebro E, Kadar L, Karlberg I, Flygare P, et al. Randomized phase II study of sequential docetaxel and irinotecan with 5-fluorouracil/folinic acid (leucovorin) in patients with advanced gastric cancer: the GATAC trial. *Gastric Cancer* 2010;**13**(3):155-61.

Guimbaud 2014 {published data only}

Guimbaud R, Bouché O, Rebischung C, Bonnetain F, Louvet C, Viret F, et al. Planned interim analysis of the intergroup FFCD-GERCOR-FNCLCC-AERO phase III study comparing two sequences of chemotherapy in locally advanced or metastatic gastric cancers. *Journal of Clinical Oncology* 2009;**210**(15 suppl):4533.

* Guimbaud R, Louvet C, Ries P, Ychou M, Maillard E, Andre T, et al. Prospective, randomized, multicenter, phase III study of fluorouracil, leucovorin, and irinotecan versus epirubicin, cisplatin, and capecitabine in advanced gastric adenocarcinoma: a French intergroup (Federation Francophone de Cancerologie Digestive, Federation Nationale des Centres de Lutte Contre le Cancer, and Groupe Cooperateur Multidisciplinaire en Oncologie) study. *Journal of Clinical Oncology* 2014;**32**:3520-6.

Gupta 1982 {published data only}

Gupta S. Treatment of advanced gastric cancer with 5-fluorouracil versus mitomycin C. *Journal of Surgical Oncology* 1982;**21**(2):94-6. [MEDLINE: 47]

Haas 1983 {published data only}

Haas C, Oishi N, Mc Donald B, Coltman C, O'Bryan R. Southwest Oncology Group Phase II-III Gastric Cancer Study: 5-fluorouracil, adriamycin, and mitomycin-C +/- vincristine (FAM vs. V-FAM) compared to chlorozotolin (CZT), M-AMSA, and dihydroxyanthracenedione (DHAD) with unimpressive differences. Proceedings/Annual Meeting of the American Society of Clinical Oncology. 1983; Vol. 2:122. [MEDLINE: 157]

Hawkins 2003 {published data only}

Hawkins R, Cunningham D, Soerbye H, Adenis A, Canon J-L, Lopez-Vivanco G, et al. Randomized phase II trial of docetaxel plus irinotecan versus docetaxel plus 5-fluorouracil (5FU) in patients with untreated advanced gastric adenocarcinoma (AGAC). Proceedings/Annual Meeting of the American Society of Clinical Oncology. 2003; Vol. 22:257. [MEDLINE: 154]

Hoffman 1998 {published data only}

Hoffman K, Glimelius B. Evaluation of clinical benefit of chemotherapy in patients with upper gastrointestinal cancer. *Acta Oncologica* 1998;**37**(7-8):651-9. [MEDLINE: 49]

Icli 1993 {published data only}

Icli F, Karaoguz H, Dincol D, Günel N. Comparison of EAP and FAM combination chemotherapies in advanced gastric cancer. Proceedings/Annual Meeting of the American Society of Clinical Oncology. 1993; Vol. 12:207. [MEDLINE: 160]

Imada 1999 {published data only}

Imada T, Sairenji M, Suda T, Yamamoto Y, Amano T, Motohashi H. A combination chemotherapy of 5-fluorouracil and cisplatin against advanced gastric cancer. *Hepato-Gastroenterology* 1999;**46**(25):594-600. [MEDLINE: 51]

Inoue 1989 {published data only}

Inoue Y, Nasu H, Nakamura J, Arakawa H, Masamune O, Yamada N, et al. Efficacy of UFT in advanced gastric carcinoma under comparative study of MMC + UFT and MMC + tegafur therapies. [Japanese]. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1989;**16**(11):3567-72. [MEDLINE: 52]

Jeung 2011 {published data only}

Jeung HC, Rha SY, Im CK, Shin SJ, Ahn JB, Yang WI, et al. A randomized phase 2 study of docetaxel and S-1 versus docetaxel and cisplatin in advanced gastric cancer with an evaluation of SPARC expression for personalized therapy. *Cancer* 2011;**117**(10):2050-7.

Kang 2007 {unpublished data only}

Kang Y, Lee J, Min Y, Lee K, Zang D, Ryoo B, et al. A randomized multi-center phase II trial of capecitabine (X) versus S-1 (S) as first-line treatment in elderly patients with metastatic or recurrent unresectable gastric cancer. *Journal of Clinical Oncology, ASCO Annual Meeting Proceedings Part 1* 2007;**25**(18S):4546.

Kelsen 1990 {published data only}

Kelsen D, Atiq O, Niedzwiecki D. A random assignment trial of fluorouracil (F), methotrexate (MTX) and adriamycin (A) (FAMTX) versus etoposide, A, and cisplatin (P) (EAP) in gastric cancer

(GAST CA). Proceedings/Annual Meeting of the American Society of Clinical Oncology. 1990; Vol. 9:121. [MEDLINE: 172]

Kilickap 2011 {published data only}

Kilickap S, Yalcin S, Ates O, Tekuzman G. The first line systemic chemotherapy in metastatic gastric carcinoma: A comparison of docetaxel, cisplatin and fluorouracil (DCF) versus cisplatin and fluorouracil (CF); versus epirubicin, cisplatin and fluorouracil (ECF) regimens in clinical setting. *Hepato-Gastroenterology* 2011;**58**(105):208-12.

Kim 1991 {published and unpublished data}

Kim NK, Park YS, Suh CI, Kang WK, Kim HT, Heo DS, et al. Phase III randomized comparison of 5-FU vs. FAM (5-FU/Adria/MMC) vs FP (5-FU/Cisplatin) in patients with advanced gastric carcinoma (AGC). Proceedings/Annual Meeting of the American Society of Clinical Oncology. 1991; Vol. 10:144. [MEDLINE: 174]

Kim 1993 {published and unpublished data}

Kim NK, Park YS, Heo DS, Suh C, Kim SY, Park KC, et al. A phase III randomized study of 5-fluorouracil and cisplatin versus 5-fluorouracil, doxorubicin, and mitomycin C versus 5-fluorouracil alone in the treatment of advanced gastric cancer. *Cancer* 1993;**71**(12):3813-8. [MEDLINE: 57]

Kim 2012 {published data only}

Kim GM, Jeung HC, Rha SY, Kim HS, Jung I, Nam BH, et al. A randomized phase II trial of S-1-oxaliplatin versus capecitabine-oxaliplatin in advanced gastric cancer. *European Journal of Cancer* 2012;**48**(4):518-26.

Kim 2013 {published data only}

Kim HS, Kim HJ, Kim SY, Lee KW, Baek SK, Kim TY, et al. Second-line chemotherapy versus supportive cancer treatment in advanced gastric cancer: A meta-analysis. *Annals of Oncology*. Oxford University Press (Great Clarendon Street, Oxford OX2 6DP, United Kingdom), 2013; Vol. 24, issue 11:2850-4.

Kitamura 1995 {published data only}

Kitamura S, Ohtani T, Kurihara M, Kosaki G, Akazawa S, Sasaki T, et al. A controlled study of AO-90, a methionine-free intravenous amino acid solution, in combination with 5-fluorouracil and mitomycin C in advanced gastric cancer patients (internal medicine group evaluation). [Japanese]. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1995;**22**(6):765-75. [MEDLINE: 58]

Koizumi 1996 {published data only}

Koizumi W, Kurihara M, Hasegawa K, Chonan A, Kubo Y, Maekawa R. Combination therapy with cisplatin, 5'-deoxy-5-fluorouridine (5'-DFUR) and mitomycin (MMC) in patients with inoperable, advanced gastric cancer: a randomized trial comparing two dosage regimens. *Oncology Reports* 1996;**3**(2):255-60. [MEDLINE: 62]

Koizumi 2004 {published data only}

Koizumi W, Fukuyama Y, Fukuda T, Akiya T, Hasegawa K, Kojima Y, et al. Randomized phase II study comparing mitomycin, cisplatin plus doxifluridine with cisplatin plus doxifluridine in advanced unresectable gastric cancer. *Anticancer Research* 2004;**24**(4):2465-70.

Koizumi 2012 {published data only}

Koizumi W, Nakayama N, Tanabe S, Sasaki T, Higuchi K, Nishimura K, et al. A multicenter phase II study of combined chemotherapy with docetaxel, cisplatin, and S-1 in patients with unresectable or recurrent gastric cancer (KDOG 0601). *Cancer Chemotherapy and Pharmacology* 2012;**69**(2):407-11.

Koizumi 2013 {published data only}

Koizumi W, Higuchi K, Shimada K, Hosaka H, Sasaki E, Nakayama N, et al. Biweekly irinotecan plus cisplatin (BIRIP) versus irinotecan alone (IRI) after S-1-based chemotherapy failure in patients with advanced gastric cancer (AGC): Final analysis of a randomised phase III trial (TCOG GI-0801/BIRIP trial). *European Journal of Cancer*. Elsevier Ltd, 2013; Vol. 49:S616.

Kolaric 1986 {published data only}

Kolaric K, Potrebica V, Stanovnik M. Controlled phase III clinical study of 4-epi-doxorubicin + 5-fluorouracil versus 5-fluorouracil alone in metastatic gastric and rectosigmoid cancer. *Oncology* 1986;**43**(2):73-7. [MEDLINE: 149]

Kondo 2000 {published data only}

Kondo K, Sakamoto J, Nakazato H, Koike A, Kitoh T, Hachisuka K, et al. A phase III randomized study comparing doxifluridine and 5-fluorouracil as supportive chemotherapy in advanced and recurrent gastric cancer. *Oncology Reports* 2000;**7**(3):485-90. [MEDLINE: 63]

Kono 2002 {published data only}

Kono K, Takahashi A, Ichihara F, Amemiya H, Iizuka H, Fujii H, et al. Prognostic significance of adoptive immunotherapy with tumor-associated lymphocytes in patients with advanced gastric cancer: a randomized trial. *Clinical Cancer Research* 2002;**8**(6):1767-71. [MEDLINE: 10]

Kornek 2002 {published data only}

Kornek GV, Raderer M, Schull B, Fiebigler W, Gedlicka C, Lenauer A, et al. Effective combination chemotherapy with paclitaxel and cisplatin with or without human granulocyte colony-stimulating factor and/or erythropoietin in patients with advanced gastric cancer. *British Journal of Cancer* 2002;**86**(12):1858-63. [MEDLINE: 8]

Kosaka 1995 {published data only}

Kosaka T, Sugaya J, Yoshida S, Takano Y, Nakano Y, Akiyama T, et al. A study of chemotherapy for patients with severely advanced gastric cancer--comparison of chemotherapy and route [Japanese]. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1995;**22**(11):1582-5. [MEDLINE: 64]

Kovach 1974 {published data only}

Kovach JS, Moertel CG, Schutt AJ, Hahn RG, Reitemeier J. A controlled study of combined 1,3 bis (2 chloroethyl) 1 nitrosourea and 5 fluorouracil therapy for advanced gastric and pancreatic cancer. *Cancer* 1974;**33**:563. [MEDLINE: 65]

Kuitunen 1991 {published data only}

Kuitunen T, Pyrhönen S. A randomized phase III trial comparing fluorouracil, epidoxorubicin and methotrexate (FEMTX) with no

treatment in nonresectable gastric cancer. *European Journal of Cancer*. 1991; Vol. 27 Supp. 2:80. [MEDLINE: 180]

Kurihara 1991 {published data only}

* Kurihara M, Izumi T, Yoshida S, Ohkubo T, Suga S, Kiyohashi A, et al. A cooperative randomized study on tegafur plus mitomycin C versus combined tegafur and uracil plus mitomycin C in the treatment of advanced gastric cancer. *Japanese Journal of Cancer Research* 1991;**82**(5):613-20.

Ohkuwa M, Ohtsu A, Boku N, Yoshida S, Miyata Y, Shirao K, et al. Long-term results for patients with unresectable gastric cancer who received chemotherapy in the Japan Clinical Oncology Group (JCOG) trials. *Gastric Cancer* 2000;**3**(3):145-50. [MEDLINE: 88]

Kurihara 1995 {published data only}

Kurihara M, Kosaki G, Taguchi T, Akazawa S, Sasaki T, Takahashi H, et al. Quality of life in patients with advanced gastric cancer receiving AO-90, a methionine-free intravenous amino acid solution, with 5-fluorouracil and mitomycin C [Japanese]. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1995;**22**(7):911-23. [MEDLINE: 67]

Kurihara 1995a {published data only}

Kurihara M, Hasegawa K, Satoh A, Koizumi W, Saigenji K, Inoue S, et al. A randomized trial investigating two cisplatin (P) dosage schedules combined 5'-DFUR (D) and mitomycin C (M) for advanced gastric cancer. Proceedings/Annual Meeting of the American Society of Clinical Oncology. 1995; Vol. 14:214. [MEDLINE: 151]

Lacave 1985 {published data only}

Lacave A, Dalesio O, Bleiberg H, Wils J, Diaz-Rubio E, Vendrik C, et al. A randomized trial of MeFA (methylccnu (Me); 5-fluorouracil (F), adriamycin (A)) vs. FA in advanced gastric cancer. Proceedings/Annual Meeting of the American Society of Clinical Oncology. 1985; Vol. 4:78. [MEDLINE: 176]

Lacave 1987 {published data only}

Lacave A, Wils J, Bleiberg H, Diaz-Rubio E, Duez N, Dalesio O. An EORTC Gastrointestinal Group phase III evaluation of combinations of methyl-CCNU, 5-fluorouracil, and adriamycin in advanced gastric cancer. *Journal of Clinical Oncology* 1987;**5**(9):1387-93. [MEDLINE: 68]

Lee 2008 {published data only}

Lee JL, Kang YK, Kang HJ, Lee KH, Zang DY, Ryoo BY, et al. A randomised multicentre phase II trial of capecitabine vs S-1 as first-line treatment in elderly patients with metastatic or recurrent unresectable gastric cancer. *British Journal of Cancer* 2008;**99**(4):584-90.

Lee 2012 {published data only}

Lee HJ, Park JC, Kim JH, Yoon JY, Cheoi KS, Lee H, et al. The ATP-based doublet chemotherapy response assay for unresectable advanced gastric cancer; a single center, prospective, randomized controlled study. *Gastrointestinal Endoscopy* 2012;**75** (4 Suppl 1):abstr. 236.

Levard 1998 {published data only}

Levard H, Pouliquen X, Hay J-M, Fingerhut A, Langlois-Zantain O, Huguier M, et al. 5-fluorouracil and cisplatin as palliative treatment of advanced oesophageal squamous cell carcinoma: a multicentre randomised controlled trial. *European Journal of Surgery* 1998;**164**:849-57. [MEDLINE: 69]

Li 2002 {published data only}

Li Q, Feng FY, Han J, Sui GJ, Zhu YG, Zhang Y, et al. Phase III clinical study of a new anticancer drug atofluding [Chinese]. *Aizheng* 2002;**21**(12):1350-3. [MEDLINE: 6]

Li 2007 {published data only}

Li XQ, Gu HG, Guo JW, Zhu XX. Clinical study of continuous venous infusion of low-dose 5-Fu and cisplatin combined with weekly docetaxel for treatment of advanced gastric cancer [Chinese]. *Modern Oncology* 2007;**15**:659-61.

Li 2011 {published data only}

Li XD, Shen H, Jiang JT, Zhang HZ, Zheng X, Shu YQ, et al. Paclitaxel based vs oxaliplatin based regimens for advanced gastric cancer. *World Journal of Gastroenterology* 2011;**17**(8):1082-7.

Li 2013 {published data only}

Li J, Qin S, Xu J, Guo W, Xiong J, Bai Y, et al. Apatinib for chemotherapy-refractory advanced metastatic gastric cancer: results from a randomized, placebo-controlled, parallel-arm, phase II trial. *Journal of Clinical Oncology* 2013;**31**(26):3219-25.

Lim 2011 {published data only}

Lim T, Yun J, Lee J, Park S, Park J, Park Y, et al. Updated survival results of the randomized phase II study comparing cisplatin/capecitabine (CX) with epirubicin plus CX (ECX) in advanced gastric cancer (AGC). *Journal of Clinical Oncology* 2011;**29** (Supp 4):Abstract 46.

Livstone 1977 {published data only}

Livstone EM. A controlled randomized evaluation of combined modality therapy (5,000 r+5-FU + MeCCNU versus combination chemotherapy (5-FU + Me CCNU) in the treatment of locally unresectable gastric carcinoma. *Digestion* 1977;**16**:256. [MEDLINE: 71]

Lordick 2013 {published data only}

Lordick F, Kang YK, Chung HC, Salman P, Oh SC, Bodoky G, et al. Capecitabine and cisplatin with or without cetuximab for patients with previously untreated advanced gastric cancer (EXPAND): a randomised, open-label phase 3 trial. *Lancet Oncology* 2013; Vol. 14, issue 6:490-9.

Lorenzen 2007 {published data only}

Lorenzen S, Hentrich M, Haberl C, Heinemann V, Schuster T, Seroneit T, et al. Split-dose docetaxel, cisplatin and leucovorin/fluorouracil as first-line therapy in advanced gastric cancer and adenocarcinoma of the gastroesophageal junction: results of a phase II trial. *Annals of Oncology* 2007;**18**(10):1673-9.

Luelmo 2006 {published data only}

Luelmo S, Polee M, Van Bochove A, Pruijt H, Ouwerkerk J, Sleeboom H, et al. Randomized phase II study of cisplatin and

high-dose 5-fluorouracil/leucovorin or paclitaxel and high-dose 5-fluorouracil/leucovorin in locally advanced or metastatic gastric cancer and adenocarcinomas of the gastroesophageal junction. *Annals of Oncology* 2006;**17**(Suppl 9: Abstract 530):ix308-ix326.

Malik 1990 {published data only}

Malik STA, Talbot D, Clarke PI, Osborne R, Reznek R, Wrigley PFM. Phase II trial of UFT in advanced colorectal and gastric cancer. *British Journal of Cancer* 1990;**62**(6):1023-5. [MEDLINE: 73]

Maruta 2007 {published data only}

Maruta F, Ishizone S, Hiraguri M, Fujimori Y, Shimizu F, Kumeda S, et al. A clinical study of docetaxel with or without 5'DFUR as a second-line chemotherapy for advanced gastric cancer. *Medical Oncology* 2007;**24**(1):71-5.

Massuti 1994 {published data only}

Massuti B, Cervantes A, Anton A, Aranda E, Diaz-Rubio E, Abad A, et al. A phase III multicenter randomized study in advanced gastric cancer (GC): fluorouracil + leucovorin + epirubicin + cisplatin (FLEP) versus fluorouracil + adriamycin + methotrexate + leucovorin (FAMTX): toxicity report. *Annals of Oncology* 1994;**5**(Suppl 8):76. [MEDLINE: 139]

Massuti 1995 {published data only}

Massuti B, Cervantes A, Aranda E, Abad A, Anton A, Jara C, et al. Myelotoxicity: the limiting side-effect of second-generation chemotherapy in gastric cancer: comparative analysis of fluorouracil + leucovorin + epirubicin + cisplatin (FLEP) and fluorouracil + adriamycin + methotrexate + leucovorin (FAMTX). Proceedings/Annual Meeting of the American Society of Clinical Oncology. 1995; Vol. 14:212. [MEDLINE: 177]

Mochiki 2012 {published data only}

Mochiki E, Ogata K, Ohno T, Toyomasu Y, Haga N, Fukai Y, et al. Phase II multi-institutional prospective randomised trial comparing S-1paclitaxel with S-1cisplatin in patients with unresectable and/or recurrent advanced gastric cancer. *British Journal of Cancer* 2012;**107**(1):31-6.

Moertel 1976 {published data only}

Moertel CG, Mittelman JA, Bakemeier RF, Engstrom P, Hanley J. Sequential and combination chemotherapy of advanced gastric cancer. *Cancer* 1976;**38**(2):678-82. [MEDLINE: 74]

Moertel 1979 {published data only}

Moertel CG, Engstrom P, Lavin PT, Gelber RD, Carbone PP. Chemotherapy of gastric and pancreatic carcinoma: a controlled evaluation of combinations of 5-fluorouracil with nitrosoureas and "lactones". *Surgery* 1979;**85**(5):509-13. [MEDLINE: 75]

Moertel 1979a {published data only}

Moertel CG, Lavin PT. Phase II-III chemotherapy studies in advanced gastric cancer. Eastern Cooperative Oncology Group. *Cancer Treatment Reports* 1979;**63**(11-12):1863-9. [MEDLINE: 76]

Moore 2005 {published data only}

Moore MJ. Three-armed trial to study new regimens for advanced gastric/esophageal cancer. *Oncology Report* 2005;**SPRING**:121-2.

Mustacchi 1997 {published data only}

Mustacchi G, Ceccherini R, Milani S, Sandri P, Leita ML, Carbonara T. Efficacy of sequential administration of G-CSF and GM-CSF after antitumor chemotherapy in patients with advanced cancer: Results of a randomized study. [Italian]. *Tumori* 1997;**83**(Suppl 5):S13-6. [MEDLINE: 78]

Nakajima 1984 {published data only}

Nakajima T, Takahashi T, Takagi K, Kuno K, Kajitani T. Comparison of 5-fluorouracil with fluorouracil in adjuvant chemotherapies with combined inductive and maintenance therapies for gastric cancer. *Journal of Clinical Oncology* 1984;**2**(12):1366-71. [MEDLINE: 79]

Nakao 1983 {published data only}

Nakao I, Uchino H, Orita K, Kaido I, Kimura T, Goto Y, et al. Clinical evaluation of schizophyllan (SPG) in advanced gastric cancer--a randomized comparative study by an envelope method. [Japanese]. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1983;**10**(4 Pt 2):1146-59. [MEDLINE: 80]

Nakashima 2008 {published data only}

Nakashima K, Hironaka S, Boku N, Onozawa Y, Fukutomi A, Yamazaki K, et al. Irinotecan plus cisplatin therapy and S-1 plus cisplatin therapy for advanced or recurrent gastric cancer in a single institution. *Japanese Journal of Clinical Oncology* 2008;**38**(12):810-5.

Niitani 1987 {published data only}

Niitani H, Kurihara M, Hasegawa K, Hatta Y, Suwa T, Tsuboi E, et al. Randomized comparison of continuous and intermittent oral administration of 5'-deoxy-5-fluorouridine in the treatment of advanced gastric cancer: a phase II trial by the multiinstitutional cooperative study group. [Japanese]. *Japanese Journal of Cancer & Chemotherapy* 1987;**14**(12):3345-50. [MEDLINE: 81]

Nordin 2001 {published data only}

Nordin K, Steel J, Hoffman K, Glimelius B. Alternative methods of interpreting quality of life data in advanced gastrointestinal cancer patients. *British Journal of Cancer* 2001;**85**(9):1265-72. [MEDLINE: 83]

Novik 1999 {published data only}

Novik Y, Ryan LM, Haller DG, Asbury R, Dutcher JP, Schutt A. Phase II protocol for the evaluation of new treatments in patients with advanced gastric carcinoma: results of ECOG 5282. *Medical Oncology* 1999;**16**(4):261-6. [MEDLINE: 84]

Ohtsu 2011 {published data only}

Ohtsu A, Shah MA, Van Cutsem E, Rha SY, Sawaki A, Park SR, et al. Bevacizumab in combination with chemotherapy as first-line therapy in advanced gastric cancer: a randomized, double-blind, placebo-controlled phase III study. *Journal of Clinical Oncology* 2011;**29**:3968-76.

Okines 2010 {published data only}

Okines AFC, Asghar U, Cunningham D, Ashley S, Ashton J, Jackson K, et al. Rechallenge with platinum plus fluoropyrimidine +/- epirubicin in patients with oesophagogastric cancer. *Oncology* 2010;**79**(1-2):150-8.

Osawa 1996 {published data only}

Osawa S, Shiroto H, Kondo Y, Nakanishi Y, Fujisawa J, Miyakawa K, et al. Randomized controlled study on adjuvant immunochemotherapy with capecitabine (HCFU) for noncuratively resected and unresected gastric cancer. [Japanese]. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1996;**23**(3):327-31. [MEDLINE: 89]

Pannettiere 1984 {published data only}

Pannettiere FJ, Haas C, McDonald B, Costanzi JJ, Talley RW, Athens J, et al. Drug combinations in the treatment of gastric adenocarcinoma: a randomized Southwest Oncology Group study. *Journal of Clinical Oncology* 1984;**2**(5):420-4. [MEDLINE: 141]

Park 2004 {published data only}

Park SH, Kim MJ, Chung M, Lee WK, Bang SM, Cho EK, et al. Interim analysis from a prospective randomized trial of taxanes plus 5-FU in advanced gastric cancer. Proceedings/Annual Meeting of the American Society of Clinical Oncology. 2004; Vol. 23:360.

Park 2006 {published data only}

Park SH, Lee WK, Chung M, Lee Y, Han SH, Bang SM, et al. Paclitaxel versus docetaxel for advanced gastric cancer: a randomized phase II trial in combination with infusional 5-fluorouracil. *Anti-Cancer Drugs* 2006;**17**(2):225-9.

Park 2008 {published data only}

Park SH, Nam E, Park J, Cho EK, Shin DB, Lee JH, et al. Randomized phase II study of irinotecan, leucovorin and 5-fluorouracil (ILF) versus cisplatin plus ILF (PILF) combination chemotherapy for advanced gastric cancer. *Annals of Oncology* 2008;**19**(4):729-33.

Popliela 1982 {published data only}

Popliela T, Zembala M, Oszacki J, Jedrychowski W. A follow-up study on chemoimmunotherapy (5-fluorouracil and BCG) in advanced gastric cancer [abstract]. *Cancer Immunology, Immunotherapy* 1982;**13**:182-4. [MEDLINE: 90]

Popov 1999 {published data only}

Popov I, Jelic S, Radosavljevic D, Nolic-Tomasevic Z. Eight-hour infusion versus bolus injection of doxorubicin in EAP regimen in patients with advanced gastric cancer (AGC): a prospective randomised trial. *European Journal of Cancer* 1999;**35**(Suppl 4):139. [MEDLINE: 178]

Popov 2000 {published and unpublished data}

Popov I, Jelic S, Radulovic S, Radosavljevic D, Nolic TZ. Eight-hour infusion versus bolus injection of doxorubicin in the EAP regimen in patients with advanced gastric cancer: a prospective randomised trial. *Annals of Oncology* 2000;**11**(3):343-8. [MEDLINE: 91]

Pozzo 2004 {published data only}

Pozzo C, Barone C, Szanto J, Padi E, Peschel C, Bukki J, et al. Irinotecan in combination with 5-fluorouracil and folinic acid or with cisplatin in patients with advanced gastric or esophageal-gastric junction adenocarcinoma: results of a randomized phase II study. *Annals of Oncology* 2004;**15**(12):1773-81.

Pyrhonen 1992 {published data only}

Pyrhonen S, Kuitunen T, Kouri M. A randomized phase III trial comparing fluorouracil, epidoxorubicin and methotrexate (FEMTX) with best supportive care in non-resectable gastric cancer. *Annals of Oncology* 1992; Vol. 3, issue Suppl 5:47. [MEDLINE: 179]

Queisser 1984 {published data only}

* Queisser W, Schnitzler G, Heim ME, König H, Katz R, Fritze D, et al. Prospective randomized study in advanced stomach cancer. Comparison between combinations of 5-fluorouracil and carmustine without and with adriamycin [Prospektiv randomisierte studie beim fortgeschrittenen magenkarzinom]. *Deutsche Medizinische Wochenschrift (1946)* 1984;**109**(25):976-80.

Schnitzler G, Queißer W, Heim ME, König HJ, Fritze D, Herrmann R, et al. Comparison of 5-FU-BCNU (FB) and 5-FU/adriamycin/BCNU (FAB) in advanced gastric cancer [Vergleich von 5-Fluorouracil-BCNU (FB) mit 5-Fluorouracil-Adriamycin-BCNU (FAB) beim fortgeschrittenen Magenkarzinom]. *Verhandlungen der Deutschen Gesellschaft für Innere Medizin* 1983;**89**:105-4.

Rake 1979 {published data only}

Rake MO, Mallinson CN, Cocking JB, Cwynarski MT, Fox CA, Wass VJ, et al. Chemotherapy in advanced gastric cancer: a controlled, prospective, randomised multi-centre study. *Gut* 1979;**20**(9):797-801. [MEDLINE: 94]

Roth 1994 {published data only}

Roth A, Kolaric K, Zupanc D. High doses of epirubicin and 5-fluorouracil with or without cisplatin in advanced gastric cancer - a preliminary report of a randomized study. *Libri Oncologici* 1994;**23**:187-93. [MEDLINE: 95]

Roth 1995 {published data only}

Roth A, Kolaric K, Zupanc D. High doses of epirubicin and 5-fluorouracil with or without cisplatin in advanced gastric cancer. *European Journal of Cancer*. 1995; Vol. 31A (Suppl 5). [MEDLINE: 182]

Roth 1997 {published data only}

Roth A, Zupanc D. Randomised clinical study (phase III) FE versus FEP in advanced gastric cancer. *European Journal of Cancer*. 1997; Vol. 33 (Suppl 8):275. [MEDLINE: 181]

Sakata 1982 {published data only}

Sakata Y, Yoshida Y, Komatsu Y, Sugawara K, Nishimura S, Kikuchi K. MQF-OK therapy in advanced terminal stomach cancer-with special reference to a comparison with MFC therapy. [Japanese]. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1982;**9**(1):109-15. [MEDLINE: 97]

Sakata 1988 {published data only}

Sakata Y, Munakata A, Baba T, Saitoh S, Itoh T, Tamura Y, et al. Controlled study of MQF-OK therapy with FT and with UFT on various advanced gastrointestinal cancers. Hirosaki Cooperative Study Group of Cancer Chemotherapy. [Japanese]. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1988;**15**(7):2065-71. [MEDLINE: 98]

Sakata 1992 {published data only}

Sakata Y, Chiba Y, Sato T, Kimura M, Fukushi G, Matsukawa M, et al. Comparative study of UFT plus mitomycin C and UFT plus doxorubicin in adenocarcinoma. Hirosaki Cooperative Group of Cancer Chemotherapy. [Japanese]. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1992;**19**(2):195-201. [MEDLINE: 99]

Sasagawa 1994 {published data only}

Sasagawa T, Ho N, Endo T, Sekine T, Sugiyama K, Tomidokoro T, et al. Randomized controlled trial of MMC + UFT and MMC + 5-FU therapy in advanced gastric cancer. [Japanese]. *Gan to Kagaku Ryoho [Japanese Journal of Cancer & Chemotherapy]* 1994;**21**(8):1179-85. [MEDLINE: 100]

Sasaki 1989 {published data only}

Sasaki T, Ota K, Ibayashi J, Sakata Y, Matsuoka T, Ishikawa M. Randomized multicenter trial of sequential methotrexate and 5-fluorouracil versus 5-fluorouracil alone in advanced gastric cancer. [Japanese]. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1989;**16**(8 Pt 1):2545-55. [MEDLINE: 102]

Sasaki 1990 {published data only}

Sasaki T, Ota K, Sakata Y, Matsuoka T, Wakui A, Akazawa S, et al. High-dose leucovorin and 5-fluorouracil in advanced gastric and colorectal cancer. High-Dose Leucovorin and 5-FU Study Group. [Japanese]. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1990;**17**(12):2361-8. [MEDLINE: 103]

Sasaki 1992 {published data only}

Sasaki T. High-dose leucovorin and 5-FU. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1992;**19**(7):954-62. [MEDLINE: 101]

Sasaki 1995 {published data only}

Sasaki T. Clinical evaluation of leucovorin and 5-fluorouracil. [Japanese]. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1995;**22**(8):1001-8. [MEDLINE: 104]

Sato 1991 {published data only}

Sato H, Wakui A, Hoshi M, Kurihara M, Yokoyama M, Shimizu H. Randomized controlled trial of induced hypertension chemotherapy (IHC) using angiotensin II human (TY-10721) in advanced gastric carcinoma (TY-10721 IHC Study Group Report). [Japanese]. *Gan to Kagaku Ryoho [Japanese Journal of Cancer & Chemotherapy]* 1991;**18**(3):451-60. [MEDLINE: 105]

Sato 1995 {published data only}

Sato H, Sugiyama K, Hoshi M, Urushiyama M, Ishizuka K. Angiotensin II (All) induced hypertension chemotherapy (IHC) for unresectable gastric cancer: with reference to resection after down staging. *World Journal of Surgery* 1995;**19**(6):836-42. [MEDLINE: 106]

Satoh 2013 {published data only}

Satoh T, Doi T, Tsuji A, Omuro Y, Miwa H, Nishina T, et al. A Japanese subgroup analysis of the lapatinib for gastric cancer (TyTAN) study. *Annals of Oncology*. Oxford University Press, 2013; Vol. 24:ix6.

Satoh 2014 {published data only}

Satoh T, Xu RH, Chung HC, Sun GP, Doi T, Xu JM, et al. Lapatinib plus paclitaxel versus paclitaxel alone in the second-line treatment of HER2-amplified advanced gastric cancer in Asian populations: TyTAN--a randomized, phase III study. *Journal of Clinical Oncology* 2014;**32**:2039-49.

Schmid 2003 {published data only}

Schmid KE, Kornek GV, Schull B, Raderer M, Lenauer A, Depisch D, et al. Second-line treatment of advanced gastric cancer with oxaliplatin plus raltitrexed. *Onkologie* 2003;**26**(3):255-8. [MEDLINE: 1]

Shen 2009 {published data only}

Shen YC, Chu JH. Observation of weekly dose of docetaxel combined with small doses of cisplatin, 5-fluorouracil continuous intravenous infusion treatment of advanced gastric cancer [Chinese]. *Journal of Basic and Clinical Oncology* 2009;**22**(4):318-20.

Shin 2007 {unpublished data only}

Shin D, Lee S, Park S, Park J, Cho E, Lee J, et al. Randomized phase II trial of irinotecan, leucovorin and 5-fluorouracil (ILF) versus cisplatin plus ILF (PILF) for advanced gastric cancer. *Journal of Clinical Oncology, ASCO Annual Meeting Proceedings Part 1* 2007;**25**(18S):4580.

Shinoda 1995 {published data only}

Shinoda M, Morise K, Kusugami K, Iwase H, Ina K, Kaneko H. Combination chemotherapy with FP versus FEP in patients with advanced gastric cancer. Research group of gastric cancer chemotherapy. [Japanese]. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1995;**22**(4):515-20. [MEDLINE: 109]

Shu 1999 {published data only}

Sun H, Qun Z, Guifang L. Clinical study of 68 advanced gastric cancer cases treated by DELF project. *Zhejiang Oncology* 1999;**5**:85-6. [MEDLINE: 110]

Shudong 1996 {published data only}

Shudong X, Dehua L, Dezong Z. ACNU and methyl-CCNU in combination chemotherapy for advanced gastric cancer: a randomized comparative study. Shanghai ACNU Collaborative Study Group. [Chinese]. *Chung-Hua Chung Liu Tsa Chih [Chinese Journal of Oncology]* 1996;**18**(1):30-3. [MEDLINE: 129]

Smith 1983 {published data only}

Smith BJ, Ashford RF, Bakowski M, Hellman K, Newton K, Phillips R, et al. A trial of high-dose 5-fluorouracil with razoxane or adriamycin in the treatment of advanced adenocarcinoma of the gastrointestinal tract. *American Journal of Clinical Oncology* 1983;**6**(4):481-4. [MEDLINE: 111]

Sun 2004 {published data only}

Sun W, Whittington R, Gallagher M, O'Dwyer P, Giantonio B, Metz J, et al. Concurrent RT with 5-FU/epirubicin and cisplatin or irinotecan for locally advanced upper GI adenocarcinoma. *Oncology (Williston Park)* 2004;**18**(14 (Suppl 14)):39-42.

Sym 2013 {published data only}

Sym SJ, Hong J, Park J, Cho EK, Lee JH, Park YH, et al. A randomized phase II study of biweekly irinotecan monotherapy or a combination of irinotecan plus 5-fluorouracil/leucovorin (mFOLFIRI) in patients with metastatic gastric adenocarcinoma refractory to or progressive after first-line chemotherapy. *Cancer Chemotherapy and Pharmacology* 2013; Vol. 71, issue 2:481-8.

Taal 1990 {published data only}

Taal BG, ten Bokkel Huinink WW, Simonetti G, Franklin H, McVie JG. A phase II trial of sequential MTx and 5-FU alternated with 4-epidoxorubicin and cisplatin in advanced gastric cancer. *Cancer Investigation* 1990;**8**(5):501-4. [MEDLINE: 112]

Taguchi 1985 {published data only}

Taguchi T, Furue H, Kimura T, Kondo T, Hattori T, Itoh I, et al. Results of phase III study of lentinan. [Japanese]. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1985;**12**(2):366-78. [MEDLINE: 113]

Takahashi 1991 {published data only}

Takahashi N. 5-FU concentration in the tissue of gastric cancer, and evaluation of cancer chemotherapy with angiotensin-II. [Japanese]. *Nihon Geka Gakkai Zasshi* 1991;**92**(7):775-84. [MEDLINE: 114]

Tebbutt 2002 {published data only}

Tebbutt NC, Norman A, Cunningham D, Iveson T, Seymour M, Hickish T, et al. A multicentre, randomised phase III trial comparing protracted venous infusion (PVI) 5-fluorouracil (5-FU) with PVI 5-FU plus mitomycin C in patients with inoperable oesophago-gastric cancer. *Annals of Oncology* 2002;**13**(10):1568-75. [MEDLINE: 115]

Tebbutt 2007 {unpublished data only}

Tebbutt N, Sourjina T, Strickland A, Van Hazel G, Ganju V, Gibbs D, et al. ATTAX: Randomised phase II study evaluating weekly docetaxel-based chemotherapy combinations in advanced esophago-gastric cancer, final results of an AGITG trial. *Journal of Clinical Oncology, ASCO Annual Meeting Proceedings Part 1* 2007;**25**(18S):4528.

Tebbutt 2010 {published data only}

Tebbutt NC, Cummins MM, Sourjina T, Strickland A, Van Hazel G, Ganju V, et al. Randomised, non-comparative phase II study of weekly docetaxel with cisplatin and 5-fluorouracil or with capecitabine in oesophagogastric cancer: the AGITG ATTAX trial. *British Journal of Cancer* 2010;**102**:475-81.

Thuss-Patience 2011 {published data only}

* Thuss-Patience PC, Kretzschmar A, Bichev D, Deist T, Hinke A, Breithaupt K, et al. Survival advantage for irinotecan versus best supportive care as second-line chemotherapy in gastric cancer--a randomised phase III study of the Arbeitsgemeinschaft

Internistische Onkologie (AIO). *European Journal of Cancer* 2011;**47**(15):2306-14.

Thuss-Patience PC, Kretzschmar A, Deist T, Hinke A, Bichev D, Lebedinzew B, et al. Irinotecan versus best supportive care (BSC) as second-line therapy in gastric cancer: a randomized phase III study of the Arbeitsgemeinschaft Internistische Onkologie (AIO). *Journal of Clinical Oncology* 2009;**27**(15s):abstr 4540.

Tsushima 1991 {published data only}

Tsushima K, Sakata Y, Suzuki H, Saitoh S, Sugimoto N, Itoh T, et al. A randomized controlled study of 5-fluorouracil/doxorubicin/mitomycin C/OK-432 (FAM-OK) therapy and 4-fluorouracil/epirubicin/mitomycin C/OK-432 (FEM-OK) therapy in advanced gastric cancer. *Journal of Japan Society for Cancer Therapy* 1991;**26**(7):1317-24. [MEDLINE: 118]

Van Cutsem 2009 {published data only}

Van Cutsem E, Kang Y, Chung H, Shen L, Sawaki L, Lordick F, et al. Efficacy results from the ToGA trial: a phase III study of trastuzumab added to standard chemotherapy (CT) in first-line human epidermal growth factor receptor 2 (HER2)-positive advanced gastric cancer (GC). *Journal of Clinical Oncology Supplement* 2009;**27**:abstr LBA4509.

Vanhoefer 2000 {published data only}

* Vanhoefer Uk, Rougier P, Wilke H, Ducreux MP, Lacave AJ, Van Cutsem E, et al. Final results of a randomized phase III trial of sequential high-dose methotrexate, fluorouracil, and doxorubicin versus etoposide, leucovorin, and fluorouracil versus infusional fluorouracil and cisplatin in advanced gastric cancer: A trial of the European Organization for Research and Treatment of Cancer Gastrointestinal Tract Cancer Cooperative Group. *Journal of Clinical Oncology* 2000;**18**(14):2648-57.

Wilke H, Wils J, Rougier P, Lacave A, Van Cutsem E, Vanhoefer U, et al. Preliminary analysis of a randomized phase III trial of FAMTX versus ELF versus cisplatin/5-FU in advanced gastric cancer (GC): a trial of the EORTC Gastrointestinal Tract Cancer Cooperative Group and the AIO (Arbeitsgemeinschaft Internistische Onkologie). *Proceedings/Annual Meeting of the American Society of Clinical Oncology*. 1995; Vol. 14:206. [MEDLINE: 184]

Vaughn 1980 {published data only}

Vaughn CB, Brady P, Chinn BJ. Combination chemotherapy in advanced gastrointestinal malignancy (with 1 color plate). *Oncology* 1980;**37**(1):57-61. [MEDLINE: 120]

Vestlev 1990 {published data only}

Vestlev PM, Pedersen H. Doxorubicin and 5-fluorouracil versus doxorubicin and oral ftorafur in the treatment of advanced gastric cancer - a phase II and III trial. *Acta Oncologica* 1990;**29**(7):945-6. [MEDLINE: 121]

Villar 1987 {published data only}

Villar A, Asensio F, Candel M, Delgado F, Garcia J, Lledo S, et al. Chemotherapy of advanced gastric carcinoma (stage IV): a randomized study of FAM versus 5-FU plus BCNU. *Chemioterapia* 1987;**6**(1):57-62. [MEDLINE: 122]

Chemotherapy for advanced gastric cancer (Review)

Voznyi 1978 {published data only}

Voznyi EK, Borisov VI, Perevodchikova NI, Babaian LA, Vaarik KhM. Comparative assessment of the effectiveness of the preparations CCNU, 5-fluorouracil, ftorafur and their combinations in inoperable stomach cancer [Russian]. *Vestnik Akademii Meditsinskikh Nauk SSSR* 1978;**9**:83-7. [MEDLINE: 123]

Wadler 2002 {published data only}

Wadler S, Brain C, Catalano P, Einzig AI, Cella D, Benson AB, III. Randomized phase II trial of either fluorouracil, parenteral hydroxyurea, interferon-alpha-2a, and filgrastim or doxorubicin/docetaxel in patients with advanced gastric cancer with quality-of-life assessment: Eastern Cooperative Oncology Group Study E6296. *Cancer Journal* 2002;**8**(3):282-6. [MEDLINE: 9]

Wakui 1983 {published data only}

Wakui A, Takahashi K, Sato Y, Sato K, Matsuoka T, Saito T. Clinical evaluation of chemoimmunotherapy for advanced gastrointestinal cancer using a combined regimen of 5-fluorouracil, adriamycin and levamisole. [Japanese]. *Gan to Kagaku Ryoho [Japanese Journal of Cancer & Chemotherapy]* 1983;**10**(2):218-26. [MEDLINE: 124]

Wakui 1983a {published data only}

Wakui A, Kikuchi K, Yokoyama M, Takahashi K, Yoshida Y, Kaito I, et al. Phase III multi-center study of levamisole (LMS)--a randomized evaluation in advanced gastrointestinal cancer, with special reference to stomach cancer. [Japanese]. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1983;**10**(7):1610-23. [MEDLINE: 125]

Wakui 1986 {published data only}

Wakui A, Kasai M, Konno K, Abe R, Kanamaru R, Takahashi K, et al. Randomized study of lentinan on patients with advanced gastric and colorectal cancer. Tohoku Lentinan Study Group. [Japanese]. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1986;**13** (4 part 1):1050-9. [MEDLINE: 126]

Wang 2007 {published data only}

Wang ZH, Chen Z, Li CZ, Zhou DG, Sheng LJ, Liu B, et al. Comparative study of continuous intravenous infusional tegafur or 5-fluorouracil combined with oxaliplatin in the treatment of advanced gastric cancer. *Chinese Journal of Cancer Prevention and Treatment* 2007;**14**(2):136-9.

Waters 1999 {published data only}

Waters JS, Norman A, Cunningham D, Scarffe JH, Webb A, Harper P, et al. Long-term survival after epirubicin, cisplatin and fluorouracil for gastric cancer: results of a randomized trial. *British Journal of Cancer* 1999;**80**(1-2):269-72. [MEDLINE: 127]

Wilke 2014 {published data only}

Wilke H, Muro K, Van Cutsem E, Oh SC, Bodoky G, Shimada Y, et al. Ramucirumab plus paclitaxel versus placebo plus paclitaxel in patients with previously treated advanced gastric or gastro-oesophageal junction adenocarcinoma (RAINBOW): a double-blind, randomised phase 3 trial. *Lancet Oncology* 2014;**15**:1224-35.

Wils 1991 {published data only}

Klein HO, Buyse M, Wils JA. Prospective randomized trial using 5-fluorouracil, adriamycin and methotrexate (FAMTX) versus FAM for treatment of advanced gastric cancer. *Onkologie* 1992;**15**(5):364-7. [MEDLINE: 61]

Wils J, Klein H, Bleiberg H, Buyse M, Wagener DTh, Diaz-Rubio E, et al. EORTC 40851: A Gastrointestinal Group (GI) randomized evaluation of the toxicity of sequential high-dose methotrexate (MTX) and 5-fluorouracil (F) combined with adriamycin (A) (FAMTX) versus F, A and mitomycin C (M) (FAM) in advanced gastric cancer. Proceedings/Annual Meeting of the American Society of Clinical Oncology. 1987; Vol. 6:73. [MEDLINE: 190]

Wils J, Klein HO, Bleiberg H, Buyse M, Wagener DJ, Conroy T, et al. EORTC 40851: Sequential high dose methotrexate (MTX) and 5-fluorouracil (F) combined with adriamycin (A) (FAMTX versus F, A, and mitomycin C (M) (FAM) in advanced gastric cancer. Proceedings/Annual Meeting of the American Society of Clinical Oncology. 1989; Vol. 8:109. [MEDLINE: 186]

Wils J, Klein HO, Bleiberg H, Buyse M, Wagener DJTh, Conroy T, et al. FAMTX (5-FU, adriamycin (A) and methotrexate (MTX)): a step ahead in the treatment of advanced gastric cancer. Proceedings/Annual Meeting of the American Society of Clinical Oncology. 1990; Vol. 9, issue 102. [MEDLINE: 175]

* Wils JA, Klein HO, Wagener DJ, Bleiberg H, Reis H, Korsten F, et al. Sequential high-dose methotrexate and fluorouracil combined with doxorubicin - a step ahead in the treatment of advanced gastric cancer: a trial of the European Organization for Research and Treatment of Cancer Gastrointestinal Tract Cooperative Group. *Journal of Clinical Oncology* 1991;**9**(5):827-31.

Wils 1994 {published data only}

Wils J, Wagener DJT, Coombes RC, Fountzilias G, Bliss JM, Law M, et al. Phase III trial of fluorouracil, methotrexate and epirubicin (FEMTX) versus FEMTX plus cisplatin (FEMTX-P) in advanced gastric cancer. *Annals of Oncology*. 1994; Vol. 5 (Suppl 8). [MEDLINE: 144]

Xu 2013 {published data only}

Xu R, Ma N, Wang F, Ma L, Chen R, Chen R, et al. Results of a randomized and controlled clinical trial evaluating the efficacy and safety of combination therapy with Endostar and S-1 combined with oxaliplatin in advanced gastric cancer. *OncoTargets and Therapy* 2013;**6**:925-9.

Yamada 1994 {published data only}

Yamada Y, Tsushima K, Sakata Y, Saito S, Ito T, Sugimoto N, et al. Sequential methotrexate/5-fluorouracil therapy with 5'-deoxy-5-fluorouridine against advanced gastric cancer: comparison between bolus injection and drip infusion of 5-fluorouracil administration. Hiroasaki Cooperative Study Group for Cancer Chemotherapy. [Japanese]. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 1994;**21**(7):1029-32. [MEDLINE: 130]

Yin 1996 {published data only}

Yin Zheng-Min, Zhang Y, Xie Zhong. Elemene plus fluorouracil in the treatment of advanced gastric cancer. [Chinese]. *Chinese Journal of Clinical Oncology* 1996;**23**(11):810-12. [MEDLINE: 134]

Yoshida 2003 {published data only}

Yoshida K, Tanabe K, Ueno H, Ohta K, Hihara J, Toge T, et al. Future prospects of personalized chemotherapy in gastric cancer patients: results of a prospective randomized pilot study. *Gastric Cancer* 2003;**6**(Suppl 1):82-9. [MEDLINE: 3]

Yoshikawa 2011 {published data only}

Yoshikawa T, Tsuburaya A, Saze Z, Aoyama T, Hasegawa S, Kanemoto A, et al. [Randomized phase II trial to compare S-1 and S-1/PSK for advanced or recurrent gastric cancer—lessons from the results]. *Gan to Kagaku Ryoho. Cancer & Chemotherapy* 2011; Vol. 38, issue 12:1909-11.

Yoshino 2007 {published data only}

Yoshino S, Oka M. Randomized phase III study of S-1 alone versus S-1 plus lentinan in advanced or recurrent gastric cancer. *Biotherapy* 2007;**21**(5):315-21.

Yun 2010 {published data only}

Yun J, Park SH, Park JO, Park YS, Ho YL, Kang WK. A randomised phase II study of combination chemotherapy with epirubicin, cisplatin and capecitabine (ECX) or cisplatin and capecitabine (CX) in advanced gastric cancer. *European Journal of Cancer* 2010;**46**:885-91.

Zhao 2009 {published data only}

Zhao F, Wang Q, Zhang JW, Hang M, Chen SB. Therapeutic evaluation of docetaxel-combined chemotherapy for advanced gastric carcinoma [Chinese]. *Acta Universitatis Medicinalis NanJing (Natural Science)* 2009;**29**:237-9.

Zironi 1992 {published data only}

Zironi S, Cocconi G, Bella M, Algeri R, Bartolucci R, De Lisi V, et al. A prospective randomized trial comparing FAM combination in advanced gastric carcinoma. *Annals of Oncology*. 1992; Vol. 3 (Suppl 5):13. [MEDLINE: 165]

References to ongoing studies
Elsaid 2005 {unpublished data only}

Elsaid AA, Elkerm Y. Final results of a randomized phase III trial of docetaxel, carboplatin and 5FU versus epirubicin, cisplatin and 5FU for locally advanced gastric cancer. *Journal of Clinical Oncology, ASCO Annual Meeting Proceedings* 2005;**23**(16S):4014.

Higuchi 2012 {published data only}

Higuchi K, Koizumi W, Yamada Y, Nishikawa K, Gotoh M. Randomized phase III study of S-1 plus oxaliplatin versus S-1 plus cisplatin for first-line treatment of advanced gastric cancer. *Journal of Clinical Oncology*. 2012; Vol. 30 (suppl. 34):abstr 60.

Kurihara {unpublished data only}

Isovorin: Phase III study. Ongoing study —.

Maiello 2011 {published data only}

Maiello E, De Vita F, Gebbia V, Lorusso S, Cinieri S, Giuliani F, et al. Epirubicin (E) in combination with cisplatin (CDDP) and capecitabine (C) versus docetaxel (D) combined with 5-fluorouracil (5-FU) by continuous infusion as front-line therapy in patients with advanced gastric cancer (AGC):

Preliminary results of a randomized phase II trial of the Gruppo Oncologico Dell'Italia Meridionale. *Journal of Clinical Oncology* 2011;**29**(suppl 1):97.

NCT01498289 {published data only}

NCT01498289. S1201: Combination chemo for patients w/advanced or metastatic esophageal, gastric, or gastroesophageal junction cancer. clinicaltrials.gov/ct2/show/NCT01498289 Date first received: 22 December 2011.

NCT01558947 {published data only}

NCT01558947. Peri-operative chemotherapy with ECX or XP in the treatment of advanced gastric cancer. clinicaltrials.gov/ct2/show/NCT01558947 Date first received: 7 March 2012.

NCT01967875 {published data only}

NCT01967875. A phase 2 trial of optimizing platinum-based chemotherapy based on ERCC1 expression as first-line treatment in patients with locally advanced or metastatic gastric cancer. clinicaltrials.gov/ct2/show/NCT01967875 Date first received: 15 October 2013.

NCT02076594 {published data only}

NCT02076594. Low-Tox vs Eox in patients with locally advanced unresectable or metastatic gastric cancer. clinicaltrials.gov/ct2/show/NCT02076594 Date first received: 20 August 2013.

NCT02114359 {published data only}

NCT02114359. Chemotherapy options for the first line chemotherapy in elderly patient with advanced gastric cancer. clinicaltrials.gov/ct2/show/NCT02114359 Date first received: 9 April 2014.

NCT02289378 {published data only}

NCT02289378. Dose-dense biweekly docetaxel, oxaliplatin and 5-fluorouracil as first-line treatment in advanced gastric cancer (DaeMon-Plus). clinicaltrials.gov/ct2/show/NCT02289378 Date first received: 7 November 2014.

NCT02289547 {published data only}

NCT02289547. Phase 3 study of Xelox followed by maintenance capecitabine in the advanced gastric cancer. clinicaltrials.gov/ct2/show/NCT02289547 Date first received: 30 October 2014.

NCT02549911 {published data only}

NCT02549911. HIPEC, intravenous chemotherapy and surgery for the treatment of advanced GC with peritoneal metastasis. clinicaltrials.gov/ct2/show/NCT02549911 Date first received: 7 September 2015.

NCT02583659 {published data only}

NCT02583659. The first-line combined chemotherapy for advanced gastric cancer: a prospective observational clinical study. clinicaltrials.gov/ct2/show/NCT02583659 Date first received: 19 October 2015.

NCT02855788 {published data only}

NCT02855788. Metronomic chemotherapy in advanced gastric cancer. clinicaltrials.gov/ct2/show/NCT02855788 Date first received: 2 August 2016.

NCT03006432 {published data only}

NCT03006432. Phase iii randomised trial to evaluate folfox with or without docetaxel (TFOX) as 1st line chemotherapy for locally advanced or metastatic oesophago-gastric carcinoma (GASTFOX). clinicaltrials.gov/ct2/show/NCT03006432 Date first received: 27 December 2016.

Tsuburaya 2012 {published data only}

Tsuburaya A, Morita S, Kodera Y, Kobayashi M, Shitara K, Yamaguchi K, et al. A randomized phase II trial to elucidate the efficacy of capecitabine plus cisplatin (XP) and S-1 plus cisplatin (SP) as a first-line treatment for advanced gastric cancer: XP ascertainment vs. SP randomized PII trial (XPartS II). *BMC Cancer* 2012;**12**:307.

Additional references
Abrams 2013

Abrams JA, Gonsalves L, Neugut AI. Diverging trends in the Incidence of reflux-related and Helicobacter pylori-related gastric cardia cancer. *Journal of Clinical Gastroenterology* 2013;**47**(4):322-7.

Ajani 2007

Ajani JA, Moiseyenko VM, Tjulandin S, Majlis A, Constenla M, Boni C, et al. Clinical benefit with docetaxel plus fluorouracil and cisplatin compared with cisplatin and fluorouracil in a phase III trial of advanced gastric or gastroesophageal cancer adenocarcinoma: the V-325 Study Group. *Journal of Clinical Oncology* 2007;**25**:3205-9.

Ajani 2007a

Ajani JA, Moiseyenko VM, Tjulandin S, Majlis A, Constenla M, Boni C, et al. Quality of life with docetaxel plus cisplatin and fluorouracil compared with cisplatin and fluorouracil from a phase III trial for advanced gastric or gastroesophageal adenocarcinoma: the V-325 Study Group. *Journal of Clinical Oncology* 2007;**25**:3210-6.

Al-Batran 2010

Al-Batran SE, Ajani JA. Impact of chemotherapy on quality of life in patients with metastatic esophagogastric cancer. *Cancer* 2010;**116**(11):2511-8.

Al-Batran 2016

Al-Batran SE, Hofheinz RD, Pauligk C, Kopp HG, Haag GM, Luley KB, et al. Histopathological regression after neoadjuvant docetaxel, oxaliplatin, fluorouracil, and leucovorin versus epirubicin, cisplatin, and fluorouracil or capecitabine in patients with resectable gastric or gastro-oesophageal junction adenocarcinoma (FLOT4-AIO): results from the phase 2 part of a multicentre, open-label, randomised phase 2/3 trial. *Lancet Oncology* 2016;**17**(12):1697-708.

Altman 2001

Altman DG. Systematic reviews of evaluations of prognostic variables. In: Egger M, et al. editor(s). *Systematic Reviews in Health Care*. 2nd Edition. London: BMJ, 2001:228-47.

Bang 2010

Bang YJ, Van Cutsem E, Feyereislova A, Chung HC, Shen L, Sawaki A, et al. Trastuzumab in combination with chemotherapy versus chemotherapy alone for treatment of HER2-positive advanced gastric or gastro-oesophageal junction cancer (ToGA): a phase 3, open-label, randomised controlled trial. *Lancet* 2010;**376**(9742):687-97.

Bang 2012

Bang YJ, Kim YW, Yang HK, Chung HC, Park YK, Lee KH, et al. Adjuvant capecitabine and oxaliplatin for gastric cancer after D2 gastrectomy (CLASSIC): a phase 3 open-label, randomised controlled trial. *Lancet*. 2012/01/10 2012; Vol. 379, issue 9813:315-21. [0140-6736]

Bonenkamp 1999

Bonenkamp JJ, Hermans J, Sasako M. Extended lymph-node dissection for gastric cancer. *New England Journal of Medicine* 1999;**340**:908-14.

Bosman 2010

Bosman FT, Carneiro F, Hruban RH, Theise ND. WHO Classification of Tumours of the Digestive System. 4th Edition. Lyon (France): IARC Press, 2010. [9283224329]

Chan 2017

Chan DY, Syn NL, Yap R, Phua JN, Soh TI, Chee CE, et al. Conversion surgery post-intraperitoneal paclitaxel and systemic chemotherapy for gastric cancer carcinomatosis peritonei. Are we ready?. *Journal of Gastrointestinal Surgery* 2017;**21**(3):425-33. [DOI: [10.1007/s11605-016-3336-3](https://doi.org/10.1007/s11605-016-3336-3); NCT01739894]

Charlson 1987

Charlson ME, Pompei P, Ales KL, MacKenzie CR. A new method of classifying prognostic comorbidity in longitudinal studies: development and validation. *Journal of Chronic Diseases* 1987;**40**(5):373-83.

Chau 2004

Chau I, Norman AR, Cunningham D, Waters JS, Oates J, Ross PJ. Multivariate prognostic factor analysis in locally advanced and metastatic esophago-gastric cancer - pooled analysis from three multicenter, randomized, controlled trials using individual patient data. *Journal of Clinical Oncology* 2004;**22**(12):2395-403.

Chen 2013

Chen XL, Chen XZ, Yang C, Liao YB, Li H, Wang L, et al. Docetaxel, cisplatin and fluorouracil (DCF) regimen compared with non-taxane-containing palliative chemotherapy for gastric carcinoma: a systematic review and meta-analysis. *PLOS One* 2013;**8**:e60320.

Correa 1996

Correa P. Helicobacter pylori and gastric cancer: state of the art. *Cancer Epidemiology, Biomarkers & Prevention* 1996;**5**:477-81.

Crew 2004

Crew CD, Neugut AI. Epidemiology of upper gastrointestinal malignancies. *Seminars in Oncology* 2004;**31**(4):450-61.

Cunningham 2006

Cunningham D, Allum WH, Stenning SP, Thompson JN, Van de Velde CJ, Nicolson M, et al. Perioperative chemotherapy versus surgery alone for resectable gastroesophageal cancer. *New England Journal of Medicine* 2006;**355**(1):11-20.

Curran 2009

Curran D, Pozzo C, Zaluski J, Dank M, Barone C, Valvere V, et al. Quality of life of palliative chemotherapy naive patients with advanced adenocarcinoma of the stomach or esophagogastric junction treated with irinotecan combined with 5-fluorouracil and folinic acid: results of a randomised phase III trial. *Quality of Life Research* 2009;**18**:853-61.

Digkila 2016

Digkila A, Wagner AD. Advanced gastric cancer: Current treatment landscape and future perspectives. *World Journal of Gastroenterology* 2016;**22**(8):2403-14.

Dixon 2016

Dixon M, Mahar AL, Helyer LK, Vasilevska-Ristovska J, Law C, Coburn NG. Prognostic factors in metastatic gastric cancer: results of a population-based, retrospective cohort study in Ontario. *Gastric Cancer* 2016;**19**(1):150-9.

Ellis 2014

Ellis LM, Bernstein DS, Voest EE, Berlin JD, Sargent D, Cortazar P, et al. American Society of Clinical Oncology perspective: Raising the bar for clinical trials by defining clinically meaningful outcomes. *Journal of Clinical Oncology* 2014;**32**:1277-80.

Eremenco 2004

Eremenco SL, Cashy J, Webster K, Ohashi Y, Locker GY, Pelletier G, et al. FACT-Gastric: a new international measure of QOL in gastric cancer. Proceedings/Annual Meeting of the American Society of Clinical Oncology. 2004; Vol. 23.

Ferrell 2017

Ferrell BR, Temel JS, Temin S, Alesi ER, Balboni TA, Basch EM, et al. Integration of palliative care into standard oncology care: American Society of Clinical Oncology clinical practice guideline update. *Journal of Clinical Oncology* 2017;**35**(1):96-112.

Ferro 2014

Ferro A, Peleteiro B, Malvezzi M, Bosetti C, Bertuccio P, Levi F, et al. Worldwide trends in gastric cancer mortality (1980-2011), with predictions to 2015, and incidence by subtype. *European Journal of Cancer* 2014;**50**(7):1330-44.

Garrido 2014

Garrido M, Fonseca PJ, Vieitez JM, Frunza M, Lacave AJ. Challenges in first line chemotherapy and targeted therapy in advanced gastric cancer. *Expert Review of Anticancer Therapy* 2014;**14**(8):887-900.

GASTRIC Group 2013

GASTRIC Group, Oba K, Paoletti X, Bang YJ, Bleiberg H, Burzykowski T, et al. Role of chemotherapy for advanced/recurrent gastric cancer: an individual-patient-data meta-analysis. *European Journal of Cancer* 2013;**49**(7):1565-77.

GBD Cancer Collaboration 2017

Global Burden of Disease Cancer Collaboration. Global, Regional, and National Cancer Incidence, Mortality, Years of Life Lost, Years Lived With Disability, and Disability-Adjusted Life-years for 32 Cancer Groups, 1990 to 2015: A Systematic Analysis for the Global Burden of Disease Study. *JAMA Oncology* 2017;**3**(4):524-48.

Higgins 2003

Higgins JPT, Thompson SG, Deeks JJ, Altman DG. Measuring inconsistency in meta-analyses. *BMJ* 2003;**327**(7414):557-60.

Higgins 2008

Higgins JPT, Green S (editors). Cochrane Handbook for Systematic Reviews of Interventions Version 5.0.2 [updated September 2009]. The Cochrane Collaboration, 2008. Available from www.cochrane-handbook.org.

Higgins 2011

Higgins JPT, Green S (editors). Cochrane Handbook for Systematic Reviews of Interventions Version 5.1.0 [updated March 2011]. The Cochrane Collaboration, 2011. Available from www.cochrane-handbook.org.

Hurwitz 2004

Hurwitz H, Fehrenbacher L, Novotny W, Cartwright T, Hainsworth J, Heim W, et al. Bevacizumab plus irinotecan, fluorouracil, and leucovorin for metastatic colorectal cancer. *New England Journal of Medicine* 2004;**350**(23):2335-42.

Ilsen 2007

Ilsen DH. Docetaxel, cisplatin and fluorouracil in gastric cancer: does the punishment fit the crime?. *Journal of Clinical Oncology* 2007;**25**(22):3188-90.

Jadad 1996

Jadad AR, Moore A, Carroll D. Assessing the quality of reports of randomized clinical trials: Is blinding necessary?. *Controlled Clinical Trials* 1996;**17**:1-12.

Kang 2012

Kang JH, Lee SI, Lim DH, Park K-W, Oh SY, Kwon H-C, et al. Salvage chemotherapy for pretreated gastric cancer: a randomized phase III trial comparing chemotherapy plus best supportive care with best supportive care alone. *Journal of Clinical Oncology* 2012; Vol. 30, issue 13:1513-8. [0732-183X]

Kang 2016

Kang Y-K, Shah MA, Ohtsu A, Van Cutsem E, Ajani JA, van der Horst T, et al. A randomized, open-label, multicenter, adaptive phase 2/3 study of trastuzumab emtansine (T-DM1) versus a taxane (TAX) in patients (pts) with previously treated HER2-positive locally advanced or metastatic gastric/gastroesophageal junction adenocarcinoma (LA/MGC/GEJC). *Journal of Clinical Oncology* 2016;**4S**:abstr 5. [<http://meetinglibrary.asco.org/record/120278/abstract>]

Kang 2017

Kang Y-K, Satoh T, Ryu M-H, Chao Y, Kato K, Chung HC, et al. Nivolumab (ONO-4538/BMS-936558) as salvage treatment after second or later-line chemotherapy for advanced gastric

or gastro-esophageal junction cancer (AGC): A double-blinded, randomized, phase III trial. *Journal of Clinical Oncology* 2017;**35**(4S):abstract 2. [http://meetinglibrary.asco.org/record/139096/abstract]

Kataoka 2017

Kataoka K, Kinoshita T, Moehler M, Mauer M, Shitara K, Wagner A D, et al. Current management of liver metastases from gastric cancer: what is common practice? New challenge of EORTC and JCOG. *Gastric Cancer* 2017;**1 February**:Epub ahead of print.

Kelley 2003

Kelley JR, Duggan JM. Gastric cancer epidemiology and risk factors. *Journal of Clinical Epidemiology* 2003;**56**:1-9.

Kripp 2014

Kripp M, Al-Batran S E, Rosowski J, Pauligk C, Homann N, Hartmann JT, et al. Quality of life of older adult patients receiving docetaxel-based chemotherapy triplets for esophagogastric adenocarcinoma: a randomized study of the Arbeitsgemeinschaft Internistische Onkologie (AIO). *Gastric Cancer* 2014;**17**:181-7.

Lauren 1965

Lauren P. The two histological main types of gastric carcinoma: an attempt at a histological classification. *Acta Pathologica Microbiologica Scandinavica* 1965;**64**:31-49.

Leichman 1991

Leichman L, Berry BT. Cisplatin therapy for adenocarcinoma of the stomach. *Seminars in Oncology* 1991;**18 (1 Suppl 3)**:25-33.

Lordick 2014a

Lordick F, Allum W, Carneiro F, Mitry E, Tabernero J, Tan P, et al. Unmet needs and challenges in gastric cancer: the way forward. *Cancer Treatment Reviews* 2014;**40**:692-700.

Lordick 2014b

Lordick F, Lorenzen S, Yamada Y, Ilson D. Optimal chemotherapy for advanced gastric cancer: is there a global consensus?. *Gastric Cancer* 2014;**17**:213-25.

MacDonald 1992

MacDonald JS, Havlin KA. Etoposide in gastric cancer. *Seminars in Oncology* 1992;**19 (6 Suppl 13)**:59-62.

MacDonald 2001a

MacDonald JS. Gastric cancer. Educational Book. ASCO, 2001:77-80.

MacDonald 2001b

MacDonald JS, Smalley SR, Benedetti J. Chemoradiotherapy after surgery compared with surgery alone for adenocarcinoma of the stomach or gastroesophageal junction. *New England Journal of Medicine* 2001;**345**:725-30.

Machin 1997

Machin D, Stenning SP, Parmar MKP. Thirty years of Medical Research Council randomized trials in solid tumors. *Journal of Clinical Oncology* 1997;**9**:100-14.

Moher 2009

Moher D, Liberati A, Tetzlaff J, Altman DG, the PRISMA Group. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. *PLOS Medicine* 2009;**6**(7):1-6.

Munoz 1968

Munoz N, Correa P, Cuello C. Histological types of gastric carcinoma in high and low risk areas. *International Journal of Cancer* 1968;**3**:809-18.

Muro 2016

Muro K, Chung H C, Shankaran V, Geva R, Catenacci D, Gupta S, et al. Pembrolizumab for patients with PD-L1-positive advanced gastric cancer (KEYNOTE-012): a multicentre, open-label, phase 1b trial. *Lancet Oncology* 2016;**17**(6):717-26.

Norman 2003

Norman GR, Sloan J, Wyrwich KW. Interpretation of changes in health-related quality of life: the remarkable universality of half a standard deviation. *Medical Care* 2003;**41**(5):582-92.

Ohkuwa 2000

Ohkuwa M, Ohtsu A, Boku N, Yoshida S, Miyata Y, Shirao K, et al. Long-term results for patients with unresectable gastric cancer who received chemotherapy in the Japan Clinical Oncology Group (JCOG) trials. *Gastric Cancer* 2000;**3**(3):145-50. [MEDLINE: 88]

Ohtsu 2013

Ohtsu A, Ajani JA, Bai YX, Bang YJ, Chung HC, Pan HM, et al. Everolimus for previously treated advanced gastric cancer: results of the randomized, double-blind, phase III GRANITE-1 study. *Journal of Clinical Oncology*. 2013/09/18 2013; Vol. 31, issue 31:3935-43. [0732-183x]

Okines 2008

Okines A, Chau I, Cunningham D. Capecitabine in gastric cancer. *Drugs of Today* 2008;**44**(8):629-40.

Pallis 2011

Pallis AG, Ring A, Fortpied C, Penninckx B, Van Nes MC, Wedding U, et al. EORTC workshop on clinical trial methodology in older individuals with a diagnosis of solid tumors. *Annals of Oncology* 2011;**22**(8):1922-6.

Paoletti 2013

Paoletti X, Oba K, Bang YJ, Bleiberg H, Boku N, Bouche O, et al. Progression-free survival as a surrogate for overall survival in advanced/recurrent gastric cancer trials: a meta-analysis. *Journal of the National Cancer Institute* 2013;**105**(21):1667-70.

Parmar 1998

Parmar MKB, Torri V, Steward L. Extracting summary statistics to perform meta-analysis of the published literature for survival endpoints. *Statistics in Medicine* 1998;**17**:2815-34.

Peleteiro 2012

Peleteiro B, La Vecchia C, Lunet N. The role of *Helicobacter pylori* infection in the web of gastric cancer causation. *European Journal of Cancer Prevention* 2012;**21**(2):118-25.

Petrelli 2013

Petrelli F, Zaniboni A, Coinu A, Cabiddu M, Ghilardi M, Sgroi G, et al. Cisplatin or not in advanced gastric cancer: a systematic review and meta-analysis. *PLOS One* 2013;**8**:e83022.

Preusser 1988

Preusser P, Achterrath W, Wilke H, Lenaz L, Fink U, Heinicke A, et al. Chemotherapy of gastric cancer. *Cancer Treatment Reviews* 1988;**15**:257-77.

Pye 2001

Pye JK, Crumplin MK, Charles J. One-year survey of carcinoma of the esophagus and stomach in Wales. *British Journal of Surgery* 2001;**88**:278-85.

RevMan [Computer program]

Nordic Cochrane Centre, The Cochrane Collaboration. Review Manager (RevMan). Version 5.3. Copenhagen: Nordic Cochrane Centre, The Cochrane Collaboration, 2014.

Rusch 2004

Rusch VW. Are cancers of the esophagus, gastroesophageal junction, and cardia one disease, two, or several?. *Seminars in Oncology* 2004;**31**(4):444-9.

Sakuramoto 2007

Sakuramoto S, Sasako M, Yamaguchi T, Kinoshita T, Fujii M, Nashimoto A, et al. ACTS-GC Group. Adjuvant chemotherapy for gastric cancer with S-1, an oral fluoropyrimidine. *New England Journal of Medicine* 2007;**357**(18):1810-20.

Sepulveda 2002

Sepulveda C, Marlin A, Yoshida T, Ulrich A. Palliative care: the World Health Organization's global perspective. *Journal of Pain and Symptom Management* 2002;**24**(2):91-6.

Sharma 2003

Sharma P. Cancer of the esophagogastric junction: epidemiology and pathogenesis. *Journal of Gastrointestinal Surgery* 2003;**6**:516-7.

Song 2016

Song H, Zhu J, Lu D. Molecular-targeted first-line therapy for advanced gastric cancer. *Cochrane Database of Systematic Reviews* 2016, Issue 7. [DOI: [10.1002/14651858.CD011461.pub2](https://doi.org/10.1002/14651858.CD011461.pub2)]

Songun 2010

Songun I, Putter H, Kranenbarg EM, Sasako M, van de Velde CJ. Surgical treatment of gastric cancer: 15-year follow-up results of the randomised nationwide Dutch D1D2 trial. *Lancet Oncology* 2010;**11**:439-49.

Swain 2015

Swain SM, Baselga J, Kim SB, Ro J, Semiglazov V, Campone M, et al. Pertuzumab, trastuzumab, and docetaxel in HER2-positive metastatic breast cancer. *New England Journal of Medicine* 2015;**372**(8):724-34.

Syn 2015

Syn NL, Yong WP, Lee SC, Goh BC. Genetic factors affecting drug disposition in Asian cancer patients. *Expert Opinion on*

Drug Metabolism & Toxicology 2015;**11**(12):1879-92. [DOI: [10.1517/17425255.2015.1108964](https://doi.org/10.1517/17425255.2015.1108964)]

Syn 2016

Syn NL, Yong WP, Goh BC, Lee SC. Evolving landscape of tumor molecular profiling for personalized cancer therapy: a comprehensive review. *Expert Opinion on Drug Metabolism & Toxicology* 2016;**12**(8):911-22. [DOI: [10.1080/17425255.2016.1196187](https://doi.org/10.1080/17425255.2016.1196187)]

Tan 2011

Tan IB, Ivanova T, Lim KH, Ong CW, Deng N, Lee J, et al. Intrinsic subtypes of gastric cancer, based on gene expression pattern, predict survival and respond differently to chemotherapy. *Gastroenterology* 2011;**141**:476-85, e1-11.

TCGA 2014

Cancer Genome Atlas Research Network. Comprehensive molecular characterization of gastric adenocarcinoma. *Nature* 2014;**513**:202-9.

Temel 2010

Temel JS, Greer JA, Muzikansky A, Gallagher ER, Admane S, Jackson VA, et al. Early palliative care for patients with metastatic non-small-cell lung cancer. *New England Journal of Medicine* 2010;**363**(8):733-42.

Thompson 2002

Thompson SG, Higgins JB. How should meta-regression analysis be undertaken and interpreted?. *Statistics in Medicine* 2002;**21**(11):1559-73.

Tierney 2007

Tierney JF, Stewart L, Ghersi D, Burdett S, Sydes MR. Practical methods for incorporating summary time-to-event data into meta-analysis. *Trials* 2007;**8**:16.

Verma 2012

Verma S, Miles D, Gianni L, Krop IE, Welslau M, Baselga J, et al. Trastuzumab emtansine for HER2-positive advanced breast cancer. *New England Journal of Medicine* 2012;**367**(19):1783-91.

Vickery 2001

Vickery CW, Blazeby JM, Conroy T, Arraras J, Sezer O, Koller M, et al. Development of an EORTC disease-specific quality of life module for use in patients with gastric cancer. *European Journal of Cancer* 2001;**37**(8):966-71.

Wadhwa 2013

Wadhwa R, Song S, Lee JS, Yao Y, Wei Q, Ajani JA. Gastric cancer-molecular and clinical dimensions. *Nature Reviews. Clinical Oncology* 2013;**10**(11):643-55.

Wildiers 2013

Wildiers H, Mauer M, Pallis A, Hurria A, Mohile SG, Luciani A, et al. End points and trial design in geriatric oncology research: a joint European organisation for research and treatment of cancer--Alliance for Clinical Trials in Oncology--International Society Of Geriatric Oncology position article. *Journal of Clinical Oncology* 2013;**31**(29):3711-8.

Wu 2001

Wu AH, Wan P, Bernstein L. A multiethnic population-based study of smoking, alcohol and body size and risk of adenocarcinomas of the stomach and esophagus (United States). *Cancer Causes & Control* 2001;**12**:721-32.

Wu-Williams 1990

Wu-Williams AH, Yu MC, Mack TM. Lifestyle, the workplace and stomach cancer by subsite in young men of Los Angeles County. *Cancer Research* 1990;**50**:2569-76.

Ychou 2011

Ychou M, Boige V, Pignon JP, Conroy T, Bouche O, Lebreton G, et al. Perioperative chemotherapy compared with surgery alone for resectable gastroesophageal adenocarcinoma: an FNCLCC and FFCD multicenter phase III trial. *Journal of Clinical Oncology*. 2011/03/30 2011; Vol. 29, issue 13:1715-21. [0732-183x]

References to other published versions of this review
Wagner 2002

Wagner AD, Grothe W, Behl S, Kleber G, Grothey A, Haerting J, et al. Chemotherapy for advanced gastric cancer. *Cochrane*

Database of Systematic Reviews 2002, Issue 4. [DOI: [10.1002/14651858.CD004064](https://doi.org/10.1002/14651858.CD004064)]

Wagner 2005

Wagner ADW, Grothe W, Behl S, Kleber G, Grothey A, Haerting J, et al. Chemotherapy for advanced gastric cancer. *Cochrane Database of Systematic Reviews* 2005, Issue 2. [DOI: [10.1002/14651858.CD004064.pub2](https://doi.org/10.1002/14651858.CD004064.pub2)]

Wagner 2006

Wagner AD, Grothe W, Haerting J, Kleber G, Grothey A, Fleig WE, et al. Chemotherapy in advanced gastric cancer: a systematic review and meta-analysis based on aggregate data. *Journal of Clinical Oncology* 2006;**24**:2903-6.

Wagner 2010

Wagner AD, Unverzagt S, Grothe W, Kleber G, Grothey A, Haerting J, et al. Chemotherapy for advanced gastric cancer. *Cochrane Database of Systematic Reviews* 2010, Issue 3. [DOI: [10.1002/14651858.CD004064.pub3](https://doi.org/10.1002/14651858.CD004064.pub3)]

* Indicates the major publication for the study

CHARACTERISTICS OF STUDIES
Characteristics of included studies [ordered by study ID]
Ajani 2005

Methods	Multicentre RCT 2 arms Quality score: D
Participants	n = 158 (68% with tumour location in the stomach; and 32 % with tumour location at the GE-junction) Median age: 57 years ECOG 2-3: 0% (Karnofsky performance status KPS was ≥70% for all patients) Metastatic disease: 95%
Interventions	DCF: docetaxel (75 mg/m ² d1) + cisplatin (75 mg/m ² d1) + FU (750 mg/m ² /d d1-5), repeated at d 21 versus DC: docetaxel (85 mg/m ²) + cisplatin (75 mg/m ²) d 1, repeated at d 21
Outcomes	Response rates Time to progression Overall survival Toxicities
Notes	This is the phase II to chose the investigational arm in Van Cutsem 2006 . A similar proportion of patients received second-line chemotherapy (DCF 39%; DC 45%).

Ajani 2005 (Continued)

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Randomisation was centralised (Aventis, Antony, France) and was stratified for centre, liver and/or peritoneal metastases, prior gastrectomy, and measurable versus assessable disease.
Allocation concealment (selection bias)	Low risk	Randomisation was centralised (Aventis, Antony, France)
Incomplete outcome data (attrition bias) efficacy	Low risk	Modified ITT (randomised and treated patients)
Incomplete outcome data (attrition bias) safety	Low risk	Modified ITT
Selective reporting (reporting bias)	Unclear risk	All treated patients were included in the safety analyses. The primary efficacy end point was initially the CR rate in the per-protocol population. However, because CRs were infrequent in this study, the IDMC based its decision regarding treatment selection on the best ORR.
Other bias	Unclear risk	Number of diffuse adenocarcinoma is lower in the DC arm (22% vs. 38%). The protocol required that the IDMC review data on at least 70 patients (minimum of 60 assessable patients) to make their decision; however, by the time mature data on 70 patients were verified, the study had accrued 158 patients
Blinded review of CT/MRI-scans?	Low risk	All pertinent imaging studies (except for those of four patients) were reviewed by an External Response Review Committee (ERRC)

Ajani 2010

Methods	Multicentre RCT 2 arms Quality score: A
Participants	n = 1053 Median age: 59 years ECOG 2-3: 0% Metastatic disease: 96%
Interventions	S-1+Cisplatin: S-1 (50 mg/m ²) in two daily doses d1-21orally + cisplatin (75 mg/m ²) repeated at d 28 versus FU+Cisplatin: fluorouracil (1000 mg/m ² /24 hrs as 120-hour infusion) + cisplatin (100 mg/m ²), repeated at d 28

Ajani 2010 (Continued)

Cisplatin was discontinued after 6 cycles; provision to continue S-1 or FU until progression of disease or unacceptable toxicities

Outcomes	Overall survival Progression-free survival Time to treatment failure Tumour response Toxicity
Notes	Second-line therapy in 31.4% of patients, most frequently with fluoropyrimidine

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Computer-generated dynamic randomisation
Allocation concealment (selection bias)	Low risk	Centralised randomisation
Incomplete outcome data (attrition bias) efficacy	Low risk	Full analysis set of all treated patients (98.8% in S-1+cisplatin arm and 94.6% in FU+cisplatin arm)
Incomplete outcome data (attrition bias) safety	Low risk	Full analysis set of all treated patients
Selective reporting (reporting bias)	Low risk	PEP: OS SEP: ORR, PFS, TTF, safety
Other bias	Low risk	None
Blinded review of CT/MRI-scans?	Low risk	Radiographic evidence of response to treatment was also independently reviewed. An independent data monitoring committee oversaw the safety and efficacy data along with other aspects of the conduct of the study.

Al Batran 2008

Methods	Multicentre RCT 2 arms Quality score: D
Participants	n = 220 Median age: 64 years ECOG 2-3: 9% Metastatic disease: 94%
Interventions	FLO: oxaliplatin 85 mg/m ² ; leucovorin 200 mg/m ² and FU 2.600 mg/m ² as 24-hour continuous infusion every 14 days

Al Batran 2008 (Continued)

FLP: cisplatin 50 mg/m²; leucovorin 200 mg/m²; FU 2.000 mg/m² weekly for 6 weeks followed by a 2-week rest

Outcomes	Median overall survival Tumour response Toxicity
Notes	A pre-planned interim analysis of toxicity and response was conducted after 80 patients were included in the study

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Unclear risk	Not stated
Incomplete outcome data (attrition bias) efficacy	Low risk	ITT analysis of all randomised patients (n = 220)
Incomplete outcome data (attrition bias) safety	Low risk	Analysis of all treated patients (n = 214)
Selective reporting (reporting bias)	Unclear risk	Report includes all expected outcomes (OS, RR and toxicity)
Other bias	High risk	Differences in baseline distribution of sex (42.9% versus 25% female) and metastatic disease (97.3% versus 90.7%). Preplanned interim analysis.
Blinded review of CT/MRI-scans?	Unclear risk	Not stated

Al-Batran 2013

Methods	Multicentre RCT 2 arms Quality score: B
Participants	n = 143 Median age: 70 years ECOG: 7.7 % Metastatic disease: 69%
Interventions	FLOT: oxaliplatin 85 mg/m ² + leucovorin 200 mg/m ² + docetaxel 50 mg/m ² , followed by 5-FU 2600 mg/m ² as a 24-hour continuous infusion d 1, repeated at 2 weeks FLO: oxaliplatin 85 mg/m ² + leucovorin 200 mg/m ² , each as infusion followed by 5-FU 2600 mg/m ² as a 24-hour continuous infusion d 1, repeated at 2 weeks until disease progression, or for a total of 8, maximum 12 cycles

Al-Batran 2013 (Continued)

Outcomes	Tolerability and feasibility, defined as per group differences in toxicity, serious adverse events, treatment duration, treatment withdrawal, discontinuation for toxicity or patient's request, proportion of patients with a ≥ 10 point change of QoL global health status (EORTC QLQ C-30 questionnaires) at eight weeks, compared to baseline
	Response rates
	Overall survival
	Progression-free survival

Notes	Study included older adult patients (age ≥ 65 years). Second line therapy was permitted (FLOT: 61% vs. FLO: 42.7%)
-------	---

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Computer random generator
Allocation concealment (selection bias)	Low risk	Centralised randomisation
Incomplete outcome data (attrition bias) efficacy	Low risk	All 143 patients (FLOT, 72; FLO, 71) were eligible for the efficacy analysis on an ITT basis
Incomplete outcome data (attrition bias) safety	Low risk	Only 1 patient was excluded from the safety analysis because of consent withdrawal before study treatment
Selective reporting (reporting bias)	Low risk	ITT analysis Expected endpoints (ORR, OS, PFS, safety by NCI-CTC etc) were included. QoL also assessed
Other bias	Low risk	The treatment arms were well balanced for pretreatment characteristics
Blinded review of CT/MRI-scans?	High risk	No blinded external radiologist

Barone 1998

Methods	Multicentre RCT 2 arms Quality score: B
Participants	n = 72 Median age: 58 years ECOG 2-3: 22%
Interventions	5-FU/Lv: Lv 100 mg/m ² ; 5-FU 370 mg/m ² d 1-5, repeated at d 29 versus EEP-L: epirubicine 30 mg/m ² d,1,5; etoposide 100 mg/m ² d 1, 3, 5; cisplatin 30 mg/m ² d 2,4 and lonidamide 150 mg/d, repeated at d 29

Chemotherapy for advanced gastric cancer (Review)

Barone 1998 (Continued)

Outcomes	Median survival 1 and 2-year survival rates Response rates Symptom control Toxicity
Notes	No standard error can be assessed for TTP

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Patients were randomised by a sealed envelope method, using random permuted blocks unknown to the clinicians, to receive either the 5-FU/6S-LV (Study A) or EEP-L combination (Study B). Patients also were stratified into four groups based on a resected or nonresected primary tumor and an ECOG PS ≤ 1 or > 1 .
Allocation concealment (selection bias)	Low risk	Sealed envelope method, using random permuted blocks unknown to the clinicians
Incomplete outcome data (attrition bias) efficacy	Low risk	One patient refused further treatment after the first cycle and was excluded from the response analysis, but not from the tolerance and survival analysis. Two patients refused chemotherapy after randomisation and were excluded completely from the analysis.
Incomplete outcome data (attrition bias) safety	Low risk	One patient refused further treatment after the first cycle and was excluded from the response analysis, but not from the tolerance and survival analysis. Two patients refused chemotherapy after randomisation and were excluded completely from the analysis.
Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	High risk	1 participant who died after the second cycle of therapy has not been included in the survival analysis = no ITT
Blinded review of CT/MRI-scans?	High risk	High risk. Likely unblinded

Boku 2009

Methods	Multicentre RCT 3 arms Quality score: A
Participants	n = 704 Median age: 63.5 years ECOG 2-3: 1.4% The study was conducted in Japan.
Interventions	5-FU: 800 mg/m ² /d, ci, d 1-5, repeated at 4 w

Boku 2009 (Continued)

versus

 IP: irinotecan 70 mg/m² d 1 +15 + cisplatin 80 mg/m² d 1, repeated at 4 w

versus

 S-1: 40 mg/m² twice a day, d 1-28, repeated at 6 w

Outcomes	Overall survival Response rates Time to treatment failure Progression-free survival Non-hospitalised survival Toxicity
Notes	This study was conducted in Japan. Aim of this study was to investigate superiority of CP and non-inferiority of S-1 to 5-FU. Second-line therapy in 78% of patients, cross-over of 39% (from 5-FU to IP), 57% (from IP to S-1) and 30% (from S-1 to IP)

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Minimisation method
Allocation concealment (selection bias)	Low risk	Central allocation
Incomplete outcome data (attrition bias) efficacy	Low risk	Extremely low rate of withdrawals
Incomplete outcome data (attrition bias) safety	Low risk	Extremely low rate of withdrawals
Selective reporting (reporting bias)	Low risk	All expected endpoints reported
Other bias	Low risk	None
Blinded review of CT/MRI-scans?	Low risk	Reviewed centrally at a trial group meeting; reviewers were unaware of treatment allocations at this time

Bouche 2004

Methods	Multicentre RCT 3 arms Quality score: D
Participants	n = 134 Median age: 65 years

Chemotherapy for advanced gastric cancer (Review)

Bouche 2004 (Continued)

Metastatic disease: 100%

Interventions	LV5FU2: LV 200 mg/m ² ; FU 400 mg/m ² bolus; FU 600 mg/m ² d 1 + 2 ,repeated at 15 d LV5FU2-cisplatin: cisplatin 50 mg/m ² d 1+ 2 + LV5FU2, repeated at d 15 LV5FU2-irinotecan: irinotecan 180 mg/m ² d 1 + LV5FU2 ,repeated at d 15
Outcomes	Tumour response Median overall survival 1 year survival rates Quality of life Toxicity
Notes	Adjuvant chemotherapy without cisplatin or irinotecan was allowed if completed at least 6 months before randomisation. Prior radiotherapy was allowed if completed more than 4 weeks before randomisation.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Low risk	Central allocation
Incomplete outcome data (attrition bias) efficacy	Low risk	One hundred thirty-six patients were enrolled between January 1999 and October 2001 in 41 centres in France. Two patients were considered ineligible; one had a lymphoma and the other had no metastatic disease. No arm was closed after the two interim analyses. Thus, the analyses were carried out on an ITT basis with the remaining 134 enrolled patients.
Incomplete outcome data (attrition bias) safety	Low risk	One hundred thirty-six patients were enrolled between January 1999 and October 2001 in 41 centres in France. Two patients were considered ineligible; one had a lymphoma and the other had no metastatic disease. No arm was closed after the two interim analyses. Thus, the analyses were carried out on an ITT basis with the remaining 134 enrolled patients.
Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	Unclear risk	Low QoL response (41% (n = 22) and 48% (n = 29) at the third evaluation in arms A and C can bias longitudinal QoL analysis. Prior chemotherapy and radiotherapy were allowed under certain circumstances.
Blinded review of CT/MRI-scans?	Unclear risk	All objective tumor responses and cases of disease stabilisation were reviewed retrospectively

Cascinu 2011

Methods	Multicentre RCT 2 arms Quality score: A
Participants	n = 78

Chemotherapy for advanced gastric cancer (Review)

Cascinu 2011 (Continued)

Median age: 63 years

ECOG 2-3: 6.3%

Metastatic disease: 89.7%

Interventions	<p>Arm A (LdCF): 5-FU (400 mg/m² bolus +600 mg/m² 22 h continuous infusion d 1-2) + cisplatin (50 mg/m² d 1) + pegylated liposomal doxorubicin (20 mg/m² d1), repeated at d 14</p> <p>versus</p> <p>Arm B (MCF) : 5-FU (400 mg/m² bolus +600 mg/m² 22 h continuous infusion d 1-2) + cisplatin (50 mg/m² d 1, repeated at d 15) + mitomycin-C (7 mg/m², repeated at d 42)</p>
Outcomes	<p>Response rates</p> <p>Time to progression</p> <p>Overall survival</p> <p>Toxicity</p>
Notes	second-line treatment in 38.5% and 25.6 % of patients

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Random sequence generated by a computer programme
Allocation concealment (selection bias)	Low risk	Central allocation
Incomplete outcome data (attrition bias) efficacy	Low risk	Seventy-seven of 78 patients were assessable for response; one patient in arm B was not assessable but was included in the ITT analysis and kept in the denominator of the response rate.
Incomplete outcome data (attrition bias) safety	Low risk	Seventy-seven of 78 patients were assessable for response; one patient in arm B was not assessable but was included in the ITT analysis and kept in the denominator of the response rate.
Selective reporting (reporting bias)	Low risk	Expected endpoints reported
Other bias	Low risk	Patients characteristics resulted well balanced between the treatment groups
Blinded review of CT/MRI-scans?	Low risk	Tumour response was assessed by an independent radiologist as central reviewer.

Chen 2015

Methods	60 AGC participants randomly divided into 2 groups by "random number table" - 30 vs 30
Participants	<p>Age range 18-75 years</p> <p>Males:Females ratio was 18:12 in the DSOX group and 14:16 in the DCF group</p>

Chen 2015 (Continued)

Interventions	Docetaxel plus S1 plus oxaliplatin (DSOX) vs Docetaxel plus fluorouracil plus cisplatin (DCF)
Outcomes	OS TTP Tumour response
Notes	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	"Random number table"
Allocation concealment (selection bias)	Unclear risk	Not stated
Incomplete outcome data (attrition bias) efficacy	Unclear risk	2 patients lost to follow-up, reasons not stated
Incomplete outcome data (attrition bias) safety	Unclear risk	2 patients lost to follow-up, reasons not stated
Selective reporting (reporting bias)	Unclear risk	Not clear how many were screened for eligibility and randomised. Only the number of evaluable patients are provided.
Other bias	Unclear risk	Nothing to comment on
Blinded review of CT/MRI-scans?	Unclear risk	Not stated

Cocconi 1994

Methods	Multicentre RCT 2 arms Quality score: A
Participants	n = 130 Metastatic disease: 88% ECOG 2-3: 7%
Interventions	FAM: 5-FU 600 mg/m ² d 1, 8, 29, 36; adriamycin 30 mg/m ² d 1, 29 and mitomycin 10 mg/m ² d 1, repeated at d 57 versus PELF: cisplatin 40 mg/m ² d 1, 5; etoposide 30 mg/m ² d 1, 5; Lv 200 mg/m ² d 1-4 + 5-FU 300 mg/m ² d 1-4, repeated at d 22
Outcomes	Median survival Response rates Time to progression Toxicity

Chemotherapy for advanced gastric cancer (Review)

Cocconi 1994 (Continued)

Notes —

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Low risk	Centralised
Incomplete outcome data (attrition bias) efficacy	Low risk	All eligible pts include in survival evaluation
Incomplete outcome data (attrition bias) safety	Low risk	All eligible pts included
Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	Low risk	Baseline characteristics well-balanced
Blinded review of CT/MRI-scans?	Low risk	Extramural review

Cocconi 2003

Methods	Multicentre RCT 2 arms Quality score: A
Participants	n = 200 Metastatic disease: 85%
Interventions	FAMTX: MTX 1500 mg/m ² d1; 5-FU 1500 mg/m ² ; Lv 7.5 mg/m ² p.o. every 6 hrs d 1-3, adriamycin 30 mg/m ² d 15, repeated at d 29 versus PELF: cisplatin 40 mg/m ² d1, 5, etoposide 30 mg/m ² d 1, 5; Lv 100 mg/m ² d 1-4, 5-FU 300 mg/m ² d 1-4, repeated at d 22
Outcomes	Median survival Response rates Time to progression Toxicity
Notes	—

Risk of bias

Bias	Authors' judgement	Support for judgement
------	--------------------	-----------------------

Cocconi 2003 (Continued)

Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Low risk	Eligible patients were centrally randomised by the operational office of GOIRC (Parma, Italy)
Incomplete outcome data (attrition bias) efficacy	Unclear risk	<p>Thirteen of the 200 randomised patients (six in the PELF and seven in the FAMTX group) did not begin the assigned chemotherapy and were not evaluated for toxicity. Reasons were not provided, and it is possible that this proportion could have an impact on analysis.</p> <p>The response of 15 patients in the PELF group and 14 in the FAMTX group were unevaluable or not evaluated for the following reasons: the treatment was never started (five versus five), protocol violations (zero versus one), insufficient treatment due to early death (five versus three), refusal (four versus two), early discontinuation due to toxicity (zero versus three) or severe medical events (one versus zero)</p>
Incomplete outcome data (attrition bias) safety	Unclear risk	Thirteen of the 200 randomised patients (six in the PELF and seven in the FAMTX group) did not begin the assigned chemotherapy and were not evaluated for toxicity. Reasons were not provided, and it is possible that this proportion could have an impact on analysis.
Selective reporting (reporting bias)	Unclear risk	Toxicity was evaluated in all of the patients receiving at least one dose of chemotherapy whether they were eligible or not
Other bias	Unclear risk	N/A
Blinded review of CT/MRI-scans?	Unclear risk	Response was assessed by the clinical investigators at each participating unit, and centrally reviewed in the case of CR, PR, no change for more than 6 months, or in the case of patients who underwent gastric resection at the end of the chemotherapy programme.

Colucci 1995

Methods	Multicentre RCT 2 arms Quality score: D
Participants	n = 71 Median age: 60 years
Interventions	5-FU/Lv: Lv 200 mg/m ² ; 5-FU 375 mg/m ² d 1-5, repeated at d 22 versus 5-FU/Lv+E: Lv 200 mg/m ² ; 5-FU 375 mg/m ² d 1-5, repeated at d 22; epirubicin 60 mg/m ² d 1, repeated at d 22
Outcomes	Median survival Response rates Secondary resectability
Notes	—

Risk of bias

Bias	Authors' judgement	Support for judgement
------	--------------------	-----------------------

Colucci 1995 (Continued)

Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Unclear risk	Not stated
Incomplete outcome data (attrition bias) efficacy	Low risk	Reasons for exclusion clearly documented and valid
Incomplete outcome data (attrition bias) safety	Low risk	Reasons for exclusion clearly documented and valid
Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	Unclear risk	No ITT, missing information about type and schedule of follow-up between groups
Blinded review of CT/MRI-scans?	High risk	Not stated, likely unblinded/by investigators

Cullinan 1985

Methods	Multicentre RCT 3 arms Quality score: D
Participants	n = 151 Metastatic disease: 62%
Interventions	5-FU included 5-FU 500 mg/m ² on days 1-5, repeated at 4 weeks, 8 weeks and every 5 weeks thereafter FAM included 5-FU at 600 mg/m ² on days 1, 8, 29 and 36; doxorubicin 30 mg/m ² on days 1 and 29, and mitomycin 10 mg/m ² on day 1 FA included 5-FU 400 mg/m ² with 40 mg of doxorubicin on day 1 every 4 weeks
Outcomes	Overall survival, response rates, toxicity (not classified according to WHO or CTC)
Notes	Study included participants with pancreatic and gastric cancer. Patients were stratified within institution according to the primary tumour. Separate results were given for participants with gastric cancer. Results only for participants with gastric cancer are included in this analysis.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Unclear risk	Not stated

Cullinan 1985 (Continued)

Incomplete outcome data (attrition bias) efficacy	Low risk	ITT analysis with 3.3% exclusion from analysis
Incomplete outcome data (attrition bias) safety	Low risk	ITT analysis
Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	High risk	5-FU/doxorubicin(FA) and FAM arm will be combined in the analysis
Blinded review of CT/MRI-scans?	High risk	High risk - likely unblinded

Cullinan 1994

Methods	Multicentre RCT 4 arms Quality score: D
Participants	n = 252 Median age: 62 years ECOG 2-3: 30%
Interventions	FAMe: 5-FU 325 mg/m ² d1-5; adriamycin 40 mg/m ² d 1, repeated at d 36; methyl-CCNU 110 mg/m ² p.o. d 1, repeated at d 71 versus FAMe+Tzt: 5-FU 325 mg/m ² , d 1-5; adriamycin 40 mg/m ² d 1, repeated at d 36; triazine 250 mg/m ² d 36-38, repeated at d 57; methyl-CCNU 110 mg/m ² p.o. d 1, repeated at d 71 versus FAP: 5-FU 300 mg/m ² d 1-5; adriamycin 40 mg/m ² d 1; cisplatin 60 mg/m ² d 1, repeated at d 36 versus FU: 5-FU 500 mg/m ² d 1-5 repeated at d 36
Outcomes	Median survival Toxicity Effects on performance status and weight gain
Notes	Three combination chemotherapy arms combined in the analysis. The single-agent 5-FU arm was opened after 56 participants were randomised. FAMe and FAP were closed after a planned interim analysis because of slightly higher death rate.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Unclear risk	Not stated
Incomplete outcome data (attrition bias)	Low risk	Low attrition rate with long follow-up (only 7 of 252 patients remain alive at the time of analysis)

Chemotherapy for advanced gastric cancer (Review)

Cullinan 1994 (Continued)
 efficacy

Incomplete outcome data (attrition bias) safety	Low risk	n = 69 + 51 + 53 + 79
Selective reporting (reporting bias)	Unclear risk	Missing response rates only a small minority of patients had measurable disease so regression rate was not used as a study endpoint
Other bias	High risk	Missing information to type and follow-up in the treatment groups, combination of 3 combination treatment arms in the analysis, 2 arms were closed after a planned interim analysis
Blinded review of CT/MRI-scans?	Unclear risk	Not stated

Dank 2008

Methods	Multicentre RCT 2 arms Quality score: D
Participants	n = 337 Median age: 59 years Metastatic disease: 95.5%
Interventions	IF: irinotecan 80 mg/m ² i.v.; FA 500 mg/m ² i.v.; 5-FU 2000 mg/m ² as a 22-hour continuous infusion on d 1 weekly for 6 weeks, followed by 1 week rest versus CF: cisplatin 100 mg/m ² i.v. d 1; 5-FU 1000 mg/m ² /day as 24-hour continuous infusion d 1-5, repeated at 4 weeks
Outcomes	Hazard ratios and median survival for overall survival and time to progression, tumour response, toxicity, QoL
Notes	The trial was planned to establish superiority or non-inferiority of IF over CF. Patients have finished prior radiotherapy and surgery 6 and 3 weeks, respectively, before randomisation. Previous adjuvant or neo-adjuvant chemotherapy was allowed if completed 12 months before first relapse.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Biased coin method
Allocation concealment (selection bias)	Unclear risk	Not stated
Incomplete outcome data (attrition bias) efficacy	Low risk	Analysis of the full-analysis population of treated patients
Incomplete outcome data (attrition bias)	Low risk	Analysis of the full-analysis population of treated patients

Chemotherapy for advanced gastric cancer (Review)

Dank 2008 (Continued)
 safety

Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	High risk	Rate of non-evaluable response was imbalanced between arms (IF 9.4% versus CF 16.8%), largely due to the higher rate of early discontinuations for toxicity in the CF arm. This difference may result from closer follow-up in IF. Prior radiotherapy and chemotherapy were allowed under certain circumstances.
Blinded review of CT/MRI-scans?	Low risk	An External Radiological Review Committee (ERRC), blinded to treatment arm, reviewed all disease assessments and determined evaluability for response and date of progression.

De Lisi 1986

Methods	Multicentre RCT 2 arms Quality score: B
Participants	n = 85 Median age: 64 years
Interventions	Arm A: 5-FU 13.5 mg/kg/day for 5 days, every 5 weeks Arm B: carmustine 50 mg/m ² on days 1 and 29; doxorubicine 25 mg/m ² on days 1, 8, 15, 29 and 36 and mitomycin C 10 mg/m ² on day 15
Outcomes	Overall survival, response rate, haematological toxicity
Notes	—

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Low risk	Centralised
Incomplete outcome data (attrition bias) efficacy	Low risk	ITT
Incomplete outcome data (attrition bias) safety	Low risk	n = 82
Selective reporting (reporting bias)	Low risk	Only haematological toxicity given (in table format - but nonhaematologic side effects were also mentioned)
Other bias	Unclear risk	N/A

De Lisi 1986 (Continued)

Blinded review of CT/MRI-scans?	High risk	High risk
---------------------------------	-----------	-----------

Dong 2014

Methods	Single-centre, randomised
Participants	<p>Total of 60 patients: 30 each received TIROX or DCF</p> <p>"Included patients: (i) gastric cancer diagnosed by pathology; (ii) patients not currently receiving chemotherapy (i.e. chemotherapy-naïve) or those who had stopped chemotherapy \geq1 month prior to enrolment"</p> <p>Enrolled "consecutive patients with recurrent or metastatic gastric cancer"</p> <p>21 and 20 pts in the TIROX and DCF group are 60 years or older</p>
Interventions	<p>"In the TIROX group, patients received 40 mg/m² S-1 orally twice daily after a meal on days 1–14; 150 mg/m² irinotecan intravenously (i.v.) infused over 90 min on the first day; 85 mg/m² oxaliplatin i.v. infused over 2 h on the first day. This treatment regimen was repeated every 21 days and a 21-day treatment period was defined as one chemotherapy cycle. In the DCF group, patients received 75 mg/m² docetaxel i.v. and 75 mg/m² cisplatin i.v. on the first day; 750 mg/m² 5-FU via continuous i.v. infusion once a day from the first day to the fifth day."</p>
Outcomes	<p>Response rates</p> <p>Safety</p>
Notes	<p>No registration number found but "All of the study methods were approved by the Ethics Committee of the First Affiliated Hospital of Zhengzhou University (no. 2010-003854)"</p>

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Low "computer-generated randomization schedule"
Allocation concealment (selection bias)	Unclear risk	None stated
Incomplete outcome data (attrition bias) efficacy	Unclear risk	Number screened for eligibility and excluded not provided. Only number of evaluable patients stated.
Incomplete outcome data (attrition bias) safety	Unclear risk	Number screened for eligibility and excluded not provided. Only number of evaluable patients stated.
Selective reporting (reporting bias)	High risk	Only response rates and certain safety data were presented. "The rates of long-term progression-free survival and overall survival were not measured". Not clear if any of these endpoints were prespecified.
Other bias	Unclear risk	N/A
Blinded review of CT/MRI-scans?	Unclear risk	Not stated

GITSG 1988

Methods	Multicentre RCT 3 arms Quality score: D
Participants	n = 249 Median age: 61 ECOG 2-3: 34%
Interventions	FAP: 5-FU 300 mg/m ² d 1,8,15,22; adriamycin 30 mg/m ² d 1; cisplatin 100 mg/m ² d 1, repeated at d 29 versus FAT: 5-FU 300 mg/m ² d1-5; adriamycin 30 mg/m ² d 1; triazinate 250 mg/m ² d 22-24, repeated at d 36 versus FAMe: 5-FU 325 mg/m ² d1-5, adriamycin 40 mg/m ² d 1, Semustine 110 mg/m ² p.os d 1, repeated at d 71
Outcomes	Median survival Response rates Toxicity not classified according to WHO or NCI
Notes	—

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Unclear risk	Not stated
Incomplete outcome data (attrition bias) efficacy	Unclear risk	Not clear how many screened for eligibility and excluded
Incomplete outcome data (attrition bias) safety	Unclear risk	Not clear how many screened for eligibility and excluded
Selective reporting (reporting bias)	Unclear risk	Toxicity not classified according to WHO or NCI
Other bias	Unclear risk	N/A
Blinded review of CT/MRI-scans?	Unclear risk	Not stated

Hironaka 2016

Methods	Multicentre, randomised, open-label, 3-armed, phase 2 trial
Participants	Median (IQR) age in the 3 arms was 65 (60–70), 65 (58–71), 65 (59–69) years
Interventions	S-1 plus leucovorin (S-1 40–60 mg orally plus oral leucovorin 25 mg twice a day for 1 week, every 2 weeks), S-1 plus leucovorin and oxaliplatin (S-1 plus leucovorin and intravenous oxaliplatin 85 mg/m ²)

Chemotherapy for advanced gastric cancer (Review)

Hironaka 2016 (Continued)

on day 1, every 2 weeks), or S-1 plus cisplatin (S-1 40–60 mg orally twice a day for 3 weeks, plus intravenous cisplatin 60 mg/m² on day 8, every 5 weeks)

49 patients were randomly assigned to the S-1 plus leucovorin group, 47 to the S-1 plus leucovorin and oxaliplatin group, and 49 to the S-1 plus cisplatin group

Outcomes Primary endpoint was overall response as assessed by an independent review committee, defined as a confirmed complete response or partial response. Secondary endpoints were overall survival, progression-free survival, time to treatment failure, disease control, duration of response, and toxic effects

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Randomisation was done centrally with the minimisation method using performance status (0 vs 1) and tumour stage (stage IV vs recurrent) as stratification factors
Allocation concealment (selection bias)	Low risk	Randomisation sequence was generated by EPS Corporation (Tokyo, Japan) independently from the study sponsor
Incomplete outcome data (attrition bias) efficacy	Low risk	After randomisation, one patient did not receive treatment because of aspiration pneumonia. †Two patients who were judged to have no measurable lesions by the independent review committee after enrolment were excluded from the efficacy analyses
Incomplete outcome data (attrition bias) safety	Low risk	After randomisation, one patient did not receive treatment because of aspiration pneumonia. †Two patients who were judged to have no measurable lesions by the independent review committee after enrolment were excluded from the efficacy analyses
Selective reporting (reporting bias)	Low risk	All prespecified outcomes were reported
Other bias	Unclear risk	N/A
Blinded review of CT/MRI-scans?	Low risk	independent data monitoring committee (but not stated review committee was blinded)

Huang 2013

Methods	Multicentre RCT 2 arms Quality score: D
Participants	n = 240 Median age: 55 years Metastatic disease: 93%
Interventions	Paclitaxel+S-1: Paclitaxel 60 mg/m ² d 1,8,15, S-1 depending on body surface area (BSA < 1.25 m ² : 80 mg/d; BSA 1.25 to <1.5 m ² : 100 mg/d; BSA > 1.5 m ² , 120 mg/d twice daily) twice daily d 1-14, repeated at d 29 versus

Chemotherapy for advanced gastric cancer (Review)

Huang 2013 (Continued)

 Paclitaxel+5-FU: Paclitaxel 60 mg/m² d 1,8,15, 5-FU 500mg/m² d 1-5, leucovorin 20 mg/m² d 1-5 repeated at d 29

Outcomes	Response rates Progression-free survival Time to treatment failure Toxicity
----------	--

Notes 6% of patients had no adenocarcinoma

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not mentioned
Allocation concealment (selection bias)	Low risk	Central randomisation
Incomplete outcome data (attrition bias) efficacy	Low risk	229/240 included in full analysis set: One patient was not eligible for the current analysis due to a lack of measurable lesions, 11 patients withdrew informed consent
Incomplete outcome data (attrition bias) safety	Low risk	229/240 included in full analysis set: One patient was not eligible for the current analysis due to a lack of measurable lesions, 11 patients withdrew informed consent
Selective reporting (reporting bias)	Low risk	All expected outcomes except OS analysed
Other bias	Unclear risk	OS not analysed
Blinded review of CT/MRI-scans?	Unclear risk	Not stated

Kang 2009

Methods	Multicentre RCT 2 arms Quality score: D
Participants	n = 316 Median age: 56 years
Interventions	XP: capecitabine 1000 mg/m ² twice daily d 1–14; cisplatin 80 mg/m ² d 1 every 3 weeks versus FP: 5-FU 800 mg/m ² /d as continuous infusion d 1–5 every 3 weeks; cisplatin 80 mg/m ² d 1 every 3 weeks
Outcomes	Progression-free-survival, overall survival Tumour response Toxicity

Kang 2009 (Continued)

Notes —

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Random permuted block design
Allocation concealment (selection bias)	Unclear risk	Not stated
Incomplete outcome data (attrition bias) efficacy	Low risk	Analysis of ITT population (n = 316)
Incomplete outcome data (attrition bias) safety	Low risk	Analysis of all treated patients (n = 311)
Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	Low risk	Patients followed up for OS till end of study regardless of withdrawal + ITT vs PP, and unadjusted vs adjusted analyses performed
Blinded review of CT/MRI-scans?	Low risk	An independent review committee (IRC) reviewed patients' radiological images and assessed tumour responses without knowledge of treatment assignment

Kikuchi 1990

Methods	Multicentre RCT 2 arms Quality score: B
Participants	n = 77 Metastatic disease: 46% ECOG 2-3: 88%
Interventions	FA: 5-FU 270 to 300 mg/m ² CI d 1- 5; adriamycin 25 mg/m ² d 5, repeated at d 22 versus FAP: 5-FU 270 to 300 mg/m ² CI d 1- 5; adriamycin 25 mg/m ² d 5; cisplatin 70 mg/m ² d 1, repeated at d 22
Outcomes	Median survival Response rates
Notes	Translated from Japanese

Risk of bias

Bias	Authors' judgement	Support for judgement
------	--------------------	-----------------------

Kikuchi 1990 (Continued)

Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Low risk	Sequential opaque envelopes
Incomplete outcome data (attrition bias) efficacy	Low risk	All evaluable pts analysed
Incomplete outcome data (attrition bias) safety	Low risk	All evaluable pts analysed
Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	Unclear risk	N/A
Blinded review of CT/MRI-scans?	High risk	Evaluated during group meetings

Kim 2001

Methods	Single-centre RCT 2 arms Quality score: B
Participants	n = 121 Metastatic disease: 90% ECOG 2-3: 12%
Interventions	FP: 5-FU 1000 mg/m ² over 6 hours d 1-5; cisplatin 60 mg/m ² d 1, repeated at d 29 versus ECF: epirubicin 50 mg/m ² d 1; cisplatin 60 mg/m ² d 1, 5-FU: 1000 mg/m ² d 1-5, repeated at d 29
Outcomes	Median survival 1- and 2-year survival rates Response rates Toxicity
Notes	Study currently published as abstract only. Final results were provided by the first author.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Abstract / supplement only
Allocation concealment (selection bias)	Unclear risk	Abstract / supplement only
Incomplete outcome data (attrition bias) efficacy	Unclear risk	Abstract / supplement only

Chemotherapy for advanced gastric cancer (Review)

Kim 2001 (Continued)

Incomplete outcome data (attrition bias) safety	Unclear risk	Abstract / supplement only
Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	Unclear risk	Abstract / supplement only
Blinded review of CT/MRI-scans?	High risk	Per first author

Kim 2014

Methods	Single-centre RCT 2 arms Quality score: D
Participants	n = 77 Median age: 57 years Metastatic disease: 62% ECOG 2-3: 13%
Interventions	wDP: docetaxel (35 mg/m ²) d 1, 8, cisplatin (60 mg/m ²) d 1 repeated at d 22 versus wDO: docetaxel (35 mg/m ²) d 1, 8, oxaliplatin (120 mg/m ²) d 1 repeated at d 22
Outcomes	Response rates Overall survival Progression-free survival Toxicity
Notes	Second-line treatment in 63 % (wDP) and 77% (wDO): irinotecan monotherapy or irinotecan plus 5-fluorouracil/ leucovorin in 67% and 70 % of the wDP and wDO arms

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Unclear risk	Not stated
Incomplete outcome data (attrition bias) efficacy	Low risk	ITT analysis

Kim 2014 (Continued)

Incomplete outcome data (attrition bias) safety	Low risk	ITT analysis
Selective reporting (reporting bias)	Low risk	All expected endpoints included
Other bias	Low risk	Patients in the wDP arm received 85% and 82 % of the planned dose intensities of docetaxel and cisplatin, respectively. In the wDO arm, the mean relative dose intensity was 83 % for docetaxel and 80 % for oxaliplatin.
Blinded review of CT/MRI-scans?	Unclear risk	Not stated

Koizumi 2008

Methods	Multicentre RCT 2 arms Quality score: A The study was conducted in Japan.
Participants	n = 305 Median age: 62 years Metastatic disease: 100% ECOG 2-3: 3%
Interventions	S-1 + cisplatin: S-1 twice daily d 1-20 repeated at d 36, dose of S-1 according to the patient's body surface area (< 1.25 m ² : 40 mg; 1.25–1.5 m ² : 50 mg; > 1.5 m ² : 60 mg) + cisplatin 60 mg/m ² d 8, repeated at d 36 versus S-1 : S-1 twice daily d 1-27 repeated at d 41, dose of S-1 according to the patient's body surface area (< 1.25 m ² : 40 mg; 1.25–1.5 m ² : 50 mg; > 1.5 m ² : 60 mg)
Outcomes	Hazard ratio for overall survival Tumour response Toxicity
Notes	This study was conducted in Japan. Due to polymorphic differences in the CYP2A6 gene in Asians and whites, the tolerability of S-1 is different in whites (Ajani 2006). The dose of S-1, which was used in this trial may not be used in non-Asian populations for this reason.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Minimisation by use of biased coin method
Allocation concealment (selection bias)	Low risk	Central allocation
Incomplete outcome data (attrition bias)	Low risk	298/305 patients included in analysis.

Koizumi 2008 (Continued)
 efficacy

Incomplete outcome data (attrition bias) safety	Low risk	298/305 patients included in analysis.
Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	Low risk	If second-line treatment was started without progressive disease (i.e. due to adverse events), patients were censored
Blinded review of CT/MRI-scans?	Low risk	Images were assessed by an extramural review committee

Koizumi 2014

Methods	Multicentre RCT 2 arms Quality score: D The study was conducted in Japan and Korea.
Participants	n = 639, 635 eligible (ITT) Median age: 65 years Advanced disease: 83%, relapse: 17% ECOG 2-3: 0%
Interventions	S-1+docetaxel: docetaxel (40mg/m ² d1) + S-1 (40-60mg/m ² -according to BSA- twice daily d 1-14), repeated at d 21 versus S-1: S-1 (40-60mg/m ² -according to BSA- twice daily d 1-28), repeated at d 42
Outcomes	Overall survival, progression-free survival, response rate, safety
Notes	The study was conducted in Japan and Korea. The dose of S-1 which was used in this trial may not be valid for non-Asian populations for this reason. This study was registered (NCT00287768).

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Unclear risk	Not stated
Incomplete outcome data (attrition bias) efficacy	Low risk	4 (0.6%) patients were ineligible (no measurable or non-measurable lesions)

Koizumi 2014 (Continued)

Incomplete outcome data (attrition bias) safety	Low risk	All treated patients were included (98%)
Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	High risk	Second-line treatment was given to 69.7% of patients in the S-1+docetaxel group and 76% in the S-1 group, planned interim analysis
Blinded review of CT/MRI-scans?	Unclear risk	Images were reviewed by a central review board.

Komatsu 2011

Methods	Multicentre RCT 2 arms Quality score: D
Participants	n = 95 Median age: 66 years ECOG 2-3: 0%
Interventions	irinotecan/S-1: irinotecan 75 mg/m ² as iv infusion d 1, 15 repeated at d 29 + S-1 initial 40–60 mg/m ² orally twice daily d 1-14, repeated at 4 weeks versus S-1: S-1 initial 40–60 mg/m ² orally twice daily d 1-28, repeated at 6 weeks In subsequent cycles doses were varied according to the most severe adverse events during the preceding cycle
Outcomes	Response rates Time to treatment failure Time to progression Overall survival Toxicity
Notes	This study was conducted in Japan.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Minimization
Allocation concealment (selection bias)	Unclear risk	Not stated
Incomplete outcome data (attrition bias)	Unclear risk	16.7 versus 9.4% were not evaluable for tumour response (RECIST)

Komatsu 2011 (Continued)
 efficacy

Incomplete outcome data (attrition bias) safety	Low risk	Two untreated patients who were excluded from safety evaluation
Selective reporting (reporting bias)	Low risk	Response rates Time to treatment failure Time to progression Overall survival Toxicity
Other bias	High risk	Patients aged over 70 years were more frequent in the group treated with irinotecan and S-1: 45.8% (irinotecan/S-1) vs. 14.9% (S-1). Median age was 70 years for patients treated with irinotecan/S-1 and 63 years for patients treated with S-1 alone.
Blinded review of CT/MRI-scans?	High risk	High risk

KRGGC 1992

Methods	Multicentre RCT 2 arms
Participants	n = 60 Inoperable or metastatic gastric adenocarcinoma
Interventions	FP: cisplatin 50 mg/m ² ; 5-FU 250 mg/m ² on day 1; 5-FU 250 mg/m ² days 2-5 FPEPIR: cisplatin 50 mg/m ² ; 5-FU 250 mg/m ² days 2-5; epirubicin 30 mg/m ² day 2
Outcomes	Response rates Overall survival Toxicity
Notes	—

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Low risk	Sealed envelopes
Incomplete outcome data (attrition bias) efficacy	High risk	5 patients not evaluated: 3 death from PD with 4 weeks of treatment, 1 due to relocation, 1 due to absence of follow-up exam

KRGGC 1992 (Continued)

Incomplete outcome data (attrition bias) safety	Unclear risk	Toxic reactions classified by "standardization of reporting of results of cancer treatment" grading
Selective reporting (re-reporting bias)	Low risk	RR, OS, toxicity (but not CTCAE/WHO)
Other bias	Unclear risk	N/A
Blinded review of CT/MRI-scans?	Unclear risk	Not stated

Levi 1986

Methods	Multicentre RCT 2 arms Quality score: B
Participants	n = 203 Median age: 60 years ECOG 2-3: 22%
Interventions	A: adriamycin 60 mg/m ² d 1, repeated at d 22 versus FAB: 5-FU 600 mg/m ² , d1,8; adriamycin 40 mg/m ² d 1, repeated at d 28 BCNU 100 mg/m ² d 1, repeated at d 56
Outcomes	Median survival 1- and 2-year survival rates Response rates Toxicity
Notes	—

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Low risk	Central
Incomplete outcome data (attrition bias) efficacy	High risk	16 patients not included because of ?inadequate follow-up, 3 = treatment cancellation, 2 = protocol violation
Incomplete outcome data (attrition bias) safety	High risk	Not ITT
Selective reporting (re-reporting bias)	Low risk	Report includes all expected outcomes

Levi 1986 (Continued)

Other bias	Unclear risk	No information about type and schedule of follow-up in the treatment groups
Blinded review of CT/MRI-scans?	Unclear risk	Not stated

Li 2014

Methods	Single-centre
Participants	n = 16 vs 16 Average age = 45.7 (range 30-65) in the FOLFOX4 group, and 42.1 (range 26-70) in the SOX group.
Interventions	"Total of 32 patients with advanced gastric cancer proved pathologically were randomly divided into 2 groups: 16 patients received SOX regimen [oxaliplatin 1.30 mg/m ² as a 2-hour infusion on day 1, S-1 capsules 80 mg/m ² ·d) twice a day per oral from day 1 to day 14 every 3 weeks], the other 16 patients received FOLFOX4 regimen [oxaliplatin 85 mg/m ² as a 2 hour infusion on day 1 and a 2 hour infusion of LV 200/(m ² ·d) followed by a 5-Fu bolus 400/(m ² ·d) and 22 hour infusion 600/(m ² ·d) for 2 consecutive days every 2 weeks]. Efficacy was evaluated at least 2 cycles"
Outcomes	Response rates Disease control rates PFS OS Safety
Notes	PFS and OS were not analysed using Kaplan-Meier methods. HR for PFS could be estimated from summary data but HR for OS could not.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Simple random assignment
Allocation concealment (selection bias)	Unclear risk	Not described
Incomplete outcome data (attrition bias) efficacy	High risk	Time-to-event analysed not analysed using Kaplan-Meier methods.
Incomplete outcome data (attrition bias) safety	High risk	Quantitative comparison only made for grade 3 or higher haematological toxicity
Selective reporting (reporting bias)	Unclear risk	N/A
Other bias	Unclear risk	N/A
Blinded review of CT/MRI-scans?	Unclear risk	Not stated

Li 2015

Methods	"prospectively recruited AGC patients all over China" (did not state number of centres involved) 2 arms: "randomly assigned in a 1:1 ratio to receive S1 plus cisplatin (CS group) or fluorouracil plus cisplatin (CF group)"
Participants	255 patients screened, 120 received at least one cycle of CS, 116 received at least one cycle of CF Mean age: CS group - 53.3 years, CF group - 55 years "About 50% of the patients had low differentiated cancer. Approximately 85% of the patients had more than one site of metastasis and over half of the patients received previous gastrectomy"
Interventions	"S-1 was given as 40mg/m ² twice daily on day 1-21 and cisplatin was 20mg/m ² iv drip on day 1-4, repeated every 5 weeks in the CS group. In the CF group, 5-Fu was given as 800 mg/m ² /d CI 120h, and the dosage of cisplatin was 20mg/m ² iv on day 1-4, repeated every 4 weeks"
Outcomes	PFS (although TTP was stated as the primary endpoint, the definition they used is more consistent with PFS) OS Safety
Notes	"As a pilot study, there is no need for sample size calculation. We planned to enroll 270 patients"

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	"Randomized grouping information for each patient was generated by central randomization system. At randomization, patients were stratified by ECOG PS (0-1 vs. 2), numbers of metastasis sites (1 vs. > 1) and gastrectomy (yes vs no)"
Allocation concealment (selection bias)	High risk	"Neither patients nor investigators were masked to treatment assignment in this open-label study"
Incomplete outcome data (attrition bias) efficacy	Unclear risk	Low rate of attrition but reasons not specified
Incomplete outcome data (attrition bias) safety	Unclear risk	Low rate of attrition but reasons not specified
Selective reporting (reporting bias)	Unclear risk	N/A
Other bias	High risk	"As a pilot study, there is no need for sample size calculation. We planned to enroll 270 patients"
Blinded review of CT/MRI-scans?	High risk	"Investigators assessed tumor response and progression"

Li 2016

Methods	Single-centre
---------	---------------

Chemotherapy for advanced gastric cancer (Review)

Li 2016 (Continued)

Non-inferiority comparison of mEOX vs FOLFIRI

RCT

2 arms: modified EOX vs FOLFIRI

Participants	105 patients (55 received EOX, 50 received FOLFIRI)
Interventions	"The EOX group was given epirubicin 50 mg/m ² iv on day one, oxaliplatin 85 mg/m ² iv on day 1 and capecitabine at a twice-daily dose of 625 mg/m ² po for 2 wk, which was repeated every 3 wk"
Outcomes	OS PFS (separately for both first- and second-line) Objective response rate Disease control rate Adverse events
Notes	Second-line chemo allowed

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not specified
Allocation concealment (selection bias)	Unclear risk	Not mentioned
Incomplete outcome data (attrition bias) efficacy	Unclear risk	Patient disposition/CONSORT flow diagram not provided; unclear how many were screened and excluded, and for what reasons
Incomplete outcome data (attrition bias) safety	Unclear risk	Patient disposition/CONSORT flow diagram not provided; unclear how many were screened and excluded, and for what reasons
Selective reporting (reporting bias)	Unclear risk	Adverse effects (NCI-CTC) categorised and analysed as grades 1-4 and 3-4
Other bias	Unclear risk	N/A
Blinded review of CT/MRI-scans?	Unclear risk	Not mentioned

Loehrer 1994

Methods	Multicentre RCT 3 arms Quality score: B
Participants	n = 165 Median age: 60 years Metastatic disease: 63%

Chemotherapy for advanced gastric cancer (Review)

Loehrer 1994 (Continued)

ECOG 2-3: 27%

Interventions	5-FU: 5-FU 500 mg/m ² d 1-5 repeated at d 29 versus E: epirubicin 90 mg/m ² d 1, repeated at d 29 versus 5-FU+E: 5-FU 400 mg/m ² d 1-5; epirubicin 90 mg/m ² d 1, repeated at d 29
Outcomes	Median survival Toxicity
Notes	—

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Random number tables
Allocation concealment (selection bias)	Low risk	Central allocation
Incomplete outcome data (attrition bias) efficacy	High risk	Analysis for all randomised and screened patients
Incomplete outcome data (attrition bias) safety	Unclear risk	Analysis for all randomised, screened and treated patients
Selective reporting (reporting bias)	High risk	No response rates are given
Other bias	Low risk	N/A
Blinded review of CT/MRI-scans?	High risk	High risk

Lu 2014

Methods	"randomized phase II clinical trial conducted at Guizhou Cancer Hospital, China"
Participants	"a total of 94 consecutive patients were enrolled to Guizhou Cancer Hospital and randomly divided into two arms: OXS group (47 cases) and S-1 group (47 cases)" In both arms, about 3/4 of participants were males Median age was 63 and 65 years in the OXS and S-1 groups
Interventions	"Advanced gastric cancer patients were treated with S-1 daily for first 2 weeks of a 3-week cycle, or S-1 daily for first 2 weeks plus 130 mg/m ² of oxaliplatin administered as a 2-hour intravenous infusion on day 1 of a 3-week cycle. S-1 was orally administered in a fixed quantity according to body surface area (BSA) as follows: BSA less than 1.25 m ² , 40 mg two times daily; 1.25,BSA,1.5 m ² , 50 mg two times daily; and BSA more than 1.5 m ² , 60 mg two times daily"

Lu 2014 (Continued)

Outcomes "The primary endpoint was OS, defined as time from date of randomization to date of death from any cause. The secondary endpoints included PFS, RR, and safety profile."

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	"Randomization was generated using a computer-generated random sequence concealed in consecutively numbered opaque sealed envelopes"
Allocation concealment (selection bias)	Low risk	"Randomization was generated using a computer-generated random sequence concealed in consecutively numbered opaque sealed envelopes"
Incomplete outcome data (attrition bias) efficacy	Low risk	No patient was lost to follow-up
Incomplete outcome data (attrition bias) safety	Low risk	No patient was lost to follow-up
Selective reporting (reporting bias)	Unclear risk	OS, PFS, RR and safety included; possible loss of information because of the way the adverse events were categorised and analysed
Other bias	Unclear risk	N/A
Blinded review of CT/MRI-scans?	Unclear risk	Not stated

Lutz 2007

Methods	Multicentre RCT 3 arms Quality score: A
Participants	n = 90 Median age: 62 years Metastatic disease: 78 % (HD-FU), 89% (HD-FU/FA); 88% (HD-FU/FA/Cis) ECOG 2-3: 8% (HD-FU), 8% (HD-FU/FA), 4% (HD-FU/FA/Cis)
Interventions	HD-FU: weekly FU 3.000 mg/m ² as 24-hour infusion versus HD-FU/FA: weeks <i>dl</i> -FA 500 mg/m ² /2 hours or <i>l</i> -FA 250 mg/m ² /2 hours + FU 2.600 mg/m ² as 24-hour infusion versus HD-FU/FA/Cis: cisplatin 50 mg/m ² /hour on days 1, 15, 29; <i>dl</i> -FA 500 mg/m ² /2 hours or <i>l</i> -FA 250 mg/m ² /2 hours; FU 2.000 mg/m ² /24-hour continuous infusion on d 1, 8, 15, 22, 29, 36. In all 3 arms, chemotherapy was administered weekly in 6 subsequent weeks, followed by 1 week rest.
Outcomes	Tumour response Median and 1-year overall survival rates Toxicity

Lutz 2007 (Continued)

Notes After stage 1 (21 patients in each arm) of the trial, the HD-FU (single agent-arm) arm was closed because only 2 responses had been observed. Total number of patients in this arm was 37 because inclusion was not interrupted before interim analysis. The results of the 2 combination arms were combined in the analysis.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Minimisation technique
Allocation concealment (selection bias)	Low risk	Central randomisation at the EORTC data centre
Incomplete outcome data (attrition bias) efficacy	Low risk	127/145 eligible, reasons for exclusions provided and valid
Incomplete outcome data (attrition bias) safety	Low risk	127/145 eligible, reasons for exclusions provided and valid
Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	High risk	Single-therapy arm was closed earlier (Simon 2-stage minimax design). The results of the 2 combination arms were combined in the analysis.
Blinded review of CT/MRI-scans?	High risk	Computed tomography scans were reviewed centrally by the study co-ordinators.

Moehler 2005

Methods	Multicentre RCT 2 arms Quality score: D
Participants	n = 120 Median age: 62 years Metastatic disease: 100%
Interventions	ILF: irinotecan 80 mg/m ² + LV 500 mg/m ² + 5-FU 2000 mg/m ² days 1, 8, 15, 22, 29, 36, repeated at 8 weeks ELF: etoposide 120 mg/m ² + LV 300 mg/m ² + 5-FU 500 mg/m ² d 1-3, repeated at d 22
Outcomes	Tumour response Median overall survival Hazard ratio
Notes	—

Risk of bias

Bias	Authors' judgement	Support for judgement
------	--------------------	-----------------------

Moehler 2005 (Continued)

Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Low risk	Central allocation by the Co-ordination Centre for Clinical trials Mainz
Incomplete outcome data (attrition bias) efficacy	Low risk	Analysis of the full-analysis set of all treated patients
Incomplete outcome data (attrition bias) safety	Low risk	Analysis of the full-analysis set of all treated patients
Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	Low risk	N/A
Blinded review of CT/MRI-scans?	Unclear risk	Not stated

Moehler 2010

Methods	Multicentre RCT 2 arms Quality score: D
Participants	n = 118 Median age: 62.5 years Metastatic disease: 100%
Interventions	XI: capecitabine 1000 mg/m ² twice daily d 1-14 + irinotecan 250 mg/m ² d 1, repeated at d 22 versus XP: capecitabine 1000 mg/m ² twice daily d 1-14 + cisplatin 80 mg/m ² d 1, repeated at d 22
Outcomes	Median overall survival 1- and 2-year rate of OS Tumour response Toxicity
Notes	The reported results are from the first stage of the study (design with adaptive interim analysis)

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Low risk	Central allocation
Incomplete outcome data (attrition bias)	Low risk	Analysis of the full-analysis set of all treated patients with at least one efficacy assessment

Moehler 2010 (Continued)
 efficacy

Incomplete outcome data (attrition bias) safety	Low risk	Analysis of the full-analysis set of all treated patients
Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	High risk	Results are from the first stage of the study (design with adaptive interim analysis)
Blinded review of CT/MRI-scans?	Unclear risk	Not stated

Murad 1993

Methods	Single-centre RCT 2 arms Quality score: not applicable
Participants	n = 41 Metastatic disease: 68% ECOG 2-3: 30%
Interventions	FAMTX: Mtx 1000 mg/m ² d1; 5-FU 1500 mg/m ² d 1; Lv 15 mg p.o. every 6 hours d 1 + 2 ,repeated at d 29 Adriamycin 30 mg/m ² d 15, repeated at d 44 versus BSC
Outcomes	Median survival Response rates Maximum toxicity for each patient Hospital admittance for toxicity
Notes	After 21 patients were randomised, further participants were directly assigned to the chemotherapy arm because of "strong evidence for benefit in the treated participants"

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Unclear risk	Not stated
Incomplete outcome data (attrition bias) efficacy	High risk	No ITT
Incomplete outcome data (attrition bias) safety	Unclear risk	No ITT

Murad 1993 (Continued)

Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	Unclear risk	After 21 participants were randomised, further participants were directly assigned to the chemotherapy arm because of "strong evidence for benefit in the treated participants"
Blinded review of CT/MRI-scans?	Unclear risk	Not stated

Narahara 2011

Methods	Multicentre RCT 2 arms Quality score: D The study was conducted in Japan.
Participants	n = 326 Median age: 63 years Metastatic disease: 20% ECOG 2-3: 3%
Interventions	S-1: oral S-1 80 mg/m ² /day d 1-28, repeated at 6 weeks S-1 + irinotecan: oral S-1 80 mg/m ² /d d 1-21 + irinotecan iv 80 mg/m ² d 1 + 15, repeated at 6 weeks
Outcomes	Overall survival Time to treatment failure Response rates Toxicity
Notes	This study was conducted in Japan. Pre-planned follow-up of ≥ 1.5 years after registration of all patients was continued to 2.5 years because of a unexpectedly high survival rate of 22% at the cut-off date after a follow-up of 1.5 years. Second-line chemotherapy was administered to 76% of patients.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Centralised dynamic allocation
Allocation concealment (selection bias)	Low risk	Centralised dynamic allocation
Incomplete outcome data (attrition bias) efficacy	Low risk	Full analysis set
Incomplete outcome data (attrition bias)	Unclear risk	4 patients found to be ineligible after starting treatment were excluded from safety analysis - ?per-protocol analysis

Narahara 2011 (Continued)
 safety

Selective reporting (reporting bias)	Low risk	Expected endpoints reported
Other bias	High risk	Second-line chemotherapy was administered to a total of 76% of patients. This is likely to dilute the effect of both treatments on overall survival, but not on progression-free-survival.
Blinded review of CT/MRI-scans?	Unclear risk	Extramural review is described (does not necessarily mean blinded)

Nishikawa 2012

Methods	Multicentre RCT 4 arms Quality score: D The study was conducted in Japan.
Participants	n = 161 Median age: 67 years ECOG 2-3: 0 %
Interventions	Group A (sequential 5-FU + paclitaxel): 5-FU 800 mg/m ² c.i.v. d 1-5, repeated at 4 weeks, followed by paclitaxel 80 mg/m ² d.i.v. d 1, 8, 15, repeated at 4 weeks after progression Group B (sequential S-1 + paclitaxel): S-1 80 mg/m ² p.o. d 1-28, repeated at 6 weeks + paclitaxel 80 mg/m ² d.i.v. d 1, 8, 15, repeated at 4 weeks after progression Group C (concomitant 5-FU + paclitaxel): 5-FU 600 mg/m ² c.i.v. d 1-5 + paclitaxel 80 mg/m ² d.i.v. d 8, 15, 22, repeated at 4 weeks Group D (concomitant S-1 + paclitaxel): S-1 80 mg/m ² p.o. d1-14 + paclitaxel 50 mg/m ² d.i.v. d1,15, repeated at 3 weeks
Outcomes	Overall survival (10 months overall survival rate) Progression-free survival Time to treatment failure Response rates Toxicity
Notes	This study was conducted in Japan. After publication of the results of the SPIRITS trial (Koizumi 2008) candidates for accrual were informed about the new treatment standard in Japan and they were offered the alternative to receive the combination therapy instead of participating in the trial.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Centralised dynamic randomisation
Allocation concealment (selection bias)	Low risk	Centralised dynamic randomisation

Chemotherapy for advanced gastric cancer (Review)

Nishikawa 2012 (Continued)

Incomplete outcome data (attrition bias) efficacy	Low risk	Only two patients in arm A and two in arm C declined therapies before the start of the assigned treatment
Incomplete outcome data (attrition bias) safety	Low risk	Only two patients in arm A and two in arm C declined therapies before the start of the assigned treatment
Selective reporting (reporting bias)	Low risk	Expected endpoints are reported
Other bias	Unclear risk	An irinotecan-containing regimen was recommended for use in case if further lines of treatment were given. No information about the percentage of patients receiving second line treatment is provided.
Blinded review of CT/MRI-scans?	Unclear risk	not stated

Ochendusko 2015

Methods	Randomised, single-centre phase 3 study	
Participants	"Most patients had metastatic disease and more than 50 % of patients in each arm have undergone gastrectomy (primary tumor resection) as part of curative or palliative treatment. Significantly more patients in the mDCF arm presented with metastases in the liver (48.1 vs. 17.2 %; p = 0.029)"	
Interventions	"The EOX regimen was given every 3 weeks, initially for a maximum of eight cycles (24 weeks of treatment). It consisted of epirubicin 50 mg/m ² (intravenous bolus), followed by oxaliplatin 130 mg/m ² (2-h intravenous infusion); capecitabine was administered orally, twice daily at the dose of 625 mg/m ² for 21 days. The mDCF regimen was administered every 2 weeks, initially for a maximum of 12 cycles (24 weeks of treatment), docetaxel 40 mg/m ² (intravenous infusion over 60 min) on day 1, followed by leucovorin 400 mg/m ² (intravenous infusion over 120 min) on day 1, followed by 5-fluorouracil 400 mg/m ² (intravenous bolus) on day 1, and then 5-fluorouracil 1000 mg/m ² /day continuous intravenous infusion on day 1 and day 2, followed by cisplatin 40 mg/m ² (intravenous infusion over 60 min) on day 3."	
Outcomes	OS	
	PFS - the definition of PFS was not clearly stated	
	Safety	

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Unclear risk	Not stated
Incomplete outcome data (attrition bias)	Low risk	No loss to follow-up

Ochendusko 2015 (Continued)

efficacy

Incomplete outcome data (attrition bias) safety	Low risk	Only one patient from mDCF arm excluded due to rapid disease progression
Selective reporting (reporting bias)	Unclear risk	RR not provided
Other bias	High risk	"Significantly more patients in the mDCF arm presented with metastases in the liver (48.1 vs. 17.2 %; p = 0.029)" - no statistical adjustment was made for baseline imbalance CT scans every 8–12 weeks; and disease progression could also be evaluated based on clinical symptoms and urgent CT was requested whenever needed
Blinded review of CT/MRI-scans?	Unclear risk	Not stated

Ocvirk 2012

Methods	Single-centre RCT 2 arms Quality score: D
Participants	n = 85 Median age: 55 years Metastatic disease: 85% ECOG 2-3: 6%
Interventions	ECF: epirubicin 50 mg/m ² i.v. d 1 + cisplatin 60 mg/m ² i.v. d 1 i.v.+ 5- FU 200 mg/m ² /day continuous infusion d 1-14, repeated at d 22 ECX: epirubicin 50 mg/m ² i.v. d1+ cisplatin 60 mg/m ² i.v. d 1 + capecitabine 825 mg/m ² orally twice daily d 1-14, repeated at d 22 Treatment was discontinued in case of disease progression, unacceptable toxicity, or if the patient refused further treatment
Outcomes	Overall survival Response rates Time to progression Toxicity
Notes	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The method of sequence generation is not described

Ocvirk 2012 (Continued)

Allocation concealment (selection bias)	Low risk	"Randomisation and allocation were done by a registration center"
Incomplete outcome data (attrition bias) efficacy	Low risk	All randomised patients were included in the ITT analysis
Incomplete outcome data (attrition bias) safety	Low risk	All randomised patients were included in the ITT analysis
Selective reporting (reporting bias)	Low risk	N/A - except that TTP rather than PFS reported
Other bias	High risk	Response assessment was done by abdominal ultrasound and/or abdominal CT (not CT of the thorax and abdomen). Both methods are insufficient.
Blinded review of CT/MRI-scans?	High risk	Response assessment was done by abdominal ultrasound and/or abdominal CT (not CT of the thorax and abdomen). Both methods are insufficient.

Ohtsu 2003

Methods	Multicentre RCT 3 arms Quality score: A
Participants	n = 280 Median age: 62 years Metastatic disease: 86% ECOG 2-3: 17%
Interventions	FU: 5-FU 800 mg/m ² CI d 1-5, repeated at d 29 versus FP: 5-FU 800 mg/m ² CI d 1-5; cisplatin 20 mg/m ² d 1-5, repeated at d 29 versus UFTM: UFT (uracil/tegafur) 375 mg/m ² twice daily p.o., mitomycin 5 mg/m ² d 1, repeated at d 8
Outcomes	Median survival Response rates Toxicity
Notes	Full information about second-line therapy given: 51% of participants in the combination therapy arms and 57% of participants in the single-agent 5-FU arm received a second-line therapy

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Minimisation
Allocation concealment (selection bias)	Low risk	JCOG data centre
Incomplete outcome data (attrition bias)	Low risk	ITT

Chemotherapy for advanced gastric cancer (Review)

Ohtsu 2003 (Continued)
 efficacy

Incomplete outcome data (attrition bias) safety	Low risk	ITT
Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	Low risk	Both combination chemotherapy arms (FP and UFTM) were combined in the analysis. High rates of second-line therapy.
Blinded review of CT/MRI-scans?	Unclear risk	"objective responses confirmed by central review at regular group meetings"

Popov 2002

Methods	Single-centre RCT 2 arms Quality score: D
Participants	n = 60 Median age: 56 years Metastatic disease: 82% ECOG 2-3: 17%
Interventions	5-FU 2600 mg/m ² over 24 hours d 1, repeated at d 15 versus EAP: etoposide 120 mg/m ² d 4-6; adriamycin 20 mg/m ² d 1,7; cisplatin 40 mg/m ² d 2, 8, repeated at d 29
Outcomes	Median survival Time to progression Response rates Toxicity
Notes	Study published as abstract, information on final results provided by first author (Popov 2002). Final publication in <i>Medical Oncology</i> 2008.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Method of random numbers
Allocation concealment (selection bias)	Unclear risk	Not stated
Incomplete outcome data (attrition bias) efficacy	Low risk	No excluded patients
Incomplete outcome data (attrition bias) safety	Low risk	No excluded patients

Popov 2002 (Continued)

Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	Unclear risk	N/A
Blinded review of CT/MRI-scans?	Low risk	Independent response review was performed by a joint interdisciplinary committee not involved in the study

Popov 2008

Methods	RCT 2 arms Quality score: D
Participants	n = 72 Median age: 56 years ECOG 2-3: 29%
Interventions	LV5-FU2 oxaliplatin: oxaliplatin 85 mg/m ² d 1+ folinic acid 200 mg/m ² , as 2-hour infusion, d 1-2 + 5-FU 400 mg/m ² , Lv. bolus d 1-2, repeated at d 15 versus LV5-FU2-CDDP: cisplatin 50 mg/m ² , d1+ folinic acid 200 mg/m ² , as 2-hour infusion, d 1-2 + 5-FU 400 mg/m ² , Lv. bolus d1-2 + 5-FU 600 mg/m ² , 22-hour continuous infusion d 1-2, repeated at d 15 The maximum number of cycles foreseen was 12
Outcomes	Median overall survival Median time to progression Tumour response rates Toxicity
Notes	—

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Unclear risk	Not stated
Incomplete outcome data (attrition bias) efficacy	High risk	Patients receiving 4 or more cycles were evaluable for efficacy
Incomplete outcome data (attrition bias) safety	Low risk	Patients receiving 1 cycle were evaluable for toxicity
Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes

Popov 2008 (Continued)

Other bias	Unclear risk	N/A
Blinded review of CT/MRI-scans?	Low risk	"Independent response review was performed by members (surgeon, medical oncologist, radiologist and pathologist) of the joint interdisciplinary committee for gastrointestinal tumors of the Institute and the University Clinic for gastrointestinal diseases. The committee members were not involved in the study"

Pyrhönen 1995

Methods	RCT 2 arms Quality score: B
Participants	n = 41 Metastatic disease: 71 %
Interventions	FEMTX: methotrexate 1500 mg/m ² d 1, 5-FU 1500 mg/m ² d 1, Lv 30 mg p.o. every 6 hours d 1, 2, epirubicin 60 mg/m ² d 15, repeated at d 29 versus BSC
Outcomes	Median survival 1-and 2-year survival rates Response rates Toxicity Palliative measures
Notes	Study terminated after 6 years when 41 participants were randomised because of slow patient accrual and "conspicuous difference in survival"

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Random permuted blocks (length 10) were used. The block was not known by clinicians.
Allocation concealment (selection bias)	Low risk	Sealed envelopes, Random permuted blocks (length 10) were used. The block was not known by clinicians.
Incomplete outcome data (attrition bias) efficacy	Low risk	Only 1 patient in treatment group did not receive at least one course of chemo
Incomplete outcome data (attrition bias) safety	Low risk	Only 1 patient in treatment group did not receive at least one course of chemo
Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	High risk	Early termination of the study

Pyrhönen 1995 (Continued)

Blinded review of CT/MRI-scans?	High risk	High risk
---------------------------------	-----------	-----------

Ridwelski 2008

Methods	Multicentre RCT 2 arms Quality score: B
Participants	n = 273 Median age: 62 years Metastatic disease: 90%
Interventions	DC: docetaxel 75 mg/m ² + cisplatin 75 mg/m ² d 1, repeated at 3 w FLC: 5-FU 2000 mg/m ² + leucovorin 500 mg/m ² d 1, 8, 15, 22, 29, 36, repeated at 7 w + cisplatin 50 mg/m ² d 1, 15, 29, repeated at 7 w (cisplatin omitted in FLC in cycle 4)
Outcomes	Median overall and hazard ratios for survival and time to progression 1-year survival Tumour response Toxicity
Notes	Study currently published as abstract only. Information on final results provided by first author.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Random number tables
Allocation concealment (selection bias)	Low risk	Central
Incomplete outcome data (attrition bias) efficacy	Low risk	Analysis for all randomised and screened patients
Incomplete outcome data (attrition bias) safety	Low risk	Analysis for all randomised, screened and treated patients
Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	Unclear risk	N/A
Blinded review of CT/MRI-scans?	High risk	Per first author

Ross 2002

Methods	Multicentre RCT
---------	-----------------

Chemotherapy for advanced gastric cancer (Review)

Ross 2002 (Continued)

 2 arms
 Quality score: B

Participants	n = 334 participants with adenocarcinoma of the stomach or gastro-oesophageal junction Median age: 55 years Metastatic disease: 68% ECOG 2-3: 18%
Interventions	ECF: epirubicin 50 mg/m ² d 1, repeated at d 22; cisplatin 60 mg/m ² d 1, repeated at d 22, 5-FU CI 300 mg/m ² continuously versus MCF: mitomycin 7 mg/m ² d 1, repeated at d 43; cisplatin 60 mg/m ² d 1, repeated at d 22, 5-FU CI 300 mg/m ² continuously
Outcomes	Median survival Response rates Quality of life Toxicity
Notes	The original study included 580 participants with inoperable adenocarcinoma, squamous cell carcinoma, or undifferentiated carcinoma of the oesophagus, oesophagogastric junction or stomach. Information on participants with gastric and gastro-oesophageal junction adenocarcinoma only was provided by the first author and is included in the meta-analysis.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Query (correspondence) to first author was not answered
Allocation concealment (selection bias)	Unclear risk	Query (correspondence) to first author was not answered
Incomplete outcome data (attrition bias) efficacy	Low risk	ITT
Incomplete outcome data (attrition bias) safety	Low risk	ITT
Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	Unclear risk	N/A
Blinded review of CT/MRI-scans?	High risk	Not done per first author correspondence

Roth 1999

Methods	Single-centre RCT 2 arms Quality score: D
---------	---

Roth 1999 (Continued)

Participants	n = 122 Median age: 55 years Metastatic disease: 67% ECOG 2-3: 48%
Interventions	FE: 5-FU 1000 mg/m ² CI d 1-5, repeated at d 29; epirubicin 120 mg/m ² d 1, repeated at d 29 versus FEP: 5-FU 1000 mg/m ² CI d 1-5, repeated at d 29, epirubicin 120 mg/m ² d 1, repeated at d 29, P 30 mg/m ² d 2, 4, repeated at d 29
Outcomes	Median survival Response rates Toxicity
Notes	—

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Unclear risk	Not stated
Incomplete outcome data (attrition bias) efficacy	Low risk	110/122 received treatment and were assessable
Incomplete outcome data (attrition bias) safety	Low risk	110/122 received treatment and were assessable
Selective reporting (reporting bias)	Low risk	All expected outcomes available
Other bias	Unclear risk	No information about type and schedule of follow-up
Blinded review of CT/MRI-scans?	High risk	High risk

Roth 2007

Methods	Multicentre RCT 3 arms Quality score: A
Participants	n = 121 Median age: 59 years Metastatic disease: 86%
Interventions	ECF: epirubicin 50 mg/m ² d 1 + cisplatin 60 mg/m ² d 1 + FU 200 mg/m ² /d as 24-hour continuous infusion on days 1-21, repeated at d 22 TC: docetaxel 85 mg/m ² d 1 + cisplatin 75 mg/m ² d 1, repeated at d 22

Chemotherapy for advanced gastric cancer (Review)

Roth 2007 (Continued)

 TCF: TC + FU 300 mg/m²/d as a 24-hour continuous infusion d 1-14, repeated at d 22 for up to 8 cycles

Outcomes	Tumour response Median overall survival and time to progression Toxicity Quality of life
Notes	Because of the toxicity of this regimen, the dose of docetaxel was reduced to 75 mg/m ² later in the trial. The results of two docetaxel arms were combined in the analysis. Second-line therapy after disease progression in 56% of patients with docetaxel and 48% of patients without docetaxel.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Low risk	Central randomisation at the SAKK Co-ordinating Center
Incomplete outcome data (attrition bias) efficacy	Low risk	"Of 121 patients randomly assigned between September 1999 and July 2003, two were not treated (renal failure, n = 1; and ineligibility, n = 1) and were excluded. Another patient received two cycles of TCF before being considered ineligible but was included in the analyses."
Incomplete outcome data (attrition bias) safety	Low risk	"Of 121 patients randomly assigned between September 1999 and July 2003, two were not treated (renal failure, n = 1; and ineligibility, n = 1) and were excluded. Another patient received two cycles of TCF before being considered ineligible but was included in the analyses."
Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	Unclear risk	N/A
Blinded review of CT/MRI-scans?	Low risk	All responses were confirmed by an independent panel of radiologists and an oncologist

Roy 2012

Methods	Multicentre RCT 2 arms Quality score: D
Participants	n = 86 Median age: 61 years Metastatic disease: 94% ECOG 2-3: 0% (Karnofsky performance status KPS was ≥70% for all patients)
Interventions	DI: docetaxel 60 mg/m ² + irinotecan 250 mg/m ² d1, repeated at d 22 DF: docetaxel 85 mg/m ² d 1 + 5-FU 750 mg/m ² d 1-5, repeated at d 22
Outcomes	Response rates Time to progression

Roy 2012 (Continued)

Time to treatment failure
 Duration of response
 Overall survival
 Toxicities
 Clinical benefit (time to definitive worsening of KPS, time to definitive weight loss, time to definitive worsening of appetite and pain-free survival)

Notes 43% patients in the DI group and 49% patients in the DF group received second-line chemotherapy (mostly a platinum containing regimen)

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Unclear risk	Not stated
Incomplete outcome data (attrition bias) efficacy	Low risk	ITT analysis with only 1 excluded because of jaundice in the DI group
Incomplete outcome data (attrition bias) safety	Low risk	ITT analysis with only 1 excluded because of jaundice in the DI group
Selective reporting (reporting bias)	Low risk	All expected endpoints included
Other bias	Unclear risk	N/A
Blinded review of CT/MRI-scans?	Unclear risk	Assessed by external response review committee

Sadighi 2006

Methods	Single-centre RCT 2 arms Quality score: D
Participants	n=86 Mean age: 56 years Metastatic disease: 100%
Interventions	ECF: epirubicin 60 mg/m ² + cisplatin 60 mg/m ² + 5-FU 750 mg/m ² /day as 5 days continuous infusion; repeated at d 22 for 6 cycles TCF: docetaxel 60 mg/m ² + cisplatin 60 mg/m ² + 5-FU 750 mg/m ² as 5 days continuous infusion; repeated at d 22 for 6 cycles
Outcomes	Response rate Quality of life

Sadighi 2006 (Continued)

 Overall survival
 Progression-free survival

Notes Extensive analysis of QoL data

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Unclear risk	Not stated
Incomplete outcome data (attrition bias) efficacy	Unclear risk	No information regarding numbers of patients beside response rates (primary endpoint)
Incomplete outcome data (attrition bias) safety	High risk	For QoL evaluation, only 71 patients were included in the comparative analysis because 15 patients did not complete the QoL measurements at the beginning of the study
Selective reporting (reporting bias)	High risk	Missing information regarding PFS as secondary endpoint
Other bias	Unclear risk	N/A
Blinded review of CT/MRI-scans?	Unclear risk	Not stated

Scheithauer 1996

Methods	Multicentre RCT 2 arms Quality score: B
Participants	n = 103 Metastatic disease: 63% ECOG 2-3: 29%
Interventions	FE: 5-FU 400 mg/m ² d 1-5; Lv 200 mg/m ² d 1-5; epirubicin 50 mg/m ² d 1, repeated at d 29 versus BSC
Outcomes	Median survival Response rates 1- and 2-year survival rates Quality of life Toxicity
Notes	Study published as abstract only; additional information provided by first author

Risk of bias

Bias	Authors' judgement	Support for judgement
------	--------------------	-----------------------

Chemotherapy for advanced gastric cancer (Review)

Scheithauer 1996 (Continued)

Random sequence generation (selection bias)	Low risk	
Allocation concealment (selection bias)	Low risk	Adequate
Incomplete outcome data (attrition bias) efficacy	Low risk	
Incomplete outcome data (attrition bias) safety	Low risk	
Selective reporting (reporting bias)	Low risk	
Other bias	Low risk	
Blinded review of CT/MRI-scans?	High risk	High risk

Shirao 2013

Methods	Multicentre RCT 2 arms Quality score: B The study was conducted in Japan .
Participants	n = 237 Median age: 60 years Metastatic disease: 100% (all patients had peritoneal metastasis), confirmed by imaging and/or ascites ECOG 2-3: 3.4%
Interventions	5-FU: 5-FU 800 mg/m ² d 1-5, repeated at d 29 versus MF: methotrexate 100 mg/m ² d 1 + 5-FU 600 mg/m ² d 1 + leucovorin rescue (leucovorin 10 mg/m ² x 6) d 1, repeated at d 8
Outcomes	Overall survival, ingestion-possible survival (=surviving days free from nutrition support in patients with sufficient ingestion at baseline, ingestive improvement in patients without sufficient ingestion at baseline), safety
Notes	The study was registered as NCT00149201 and UMIN-CTR number C000000123.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Minimization
Allocation concealment (selection bias)	Low risk	Algorithm concealed to investigators

Shirao 2013 (Continued)

Incomplete outcome data (attrition bias) efficacy	Low risk	2 non-treated patients were excluded in each group.
Incomplete outcome data (attrition bias) safety	Low risk	2 non-treated patients were excluded in each group.
Selective reporting (reporting bias)	High risk	No response and PFS reported
Other bias	High risk	Second-line treatment was given to 80.7% of patients in the 5-FU group and 72.9 in the MF group, planned interim analysis
Blinded review of CT/MRI-scans?	Unclear risk	N/A (no response and PFS reported)

Sugimoto 2014

Methods	Multicentre, randomised phase 2 "Before randomization, patients were stratified according to unresectable advanced cancer/recurrent cancer with adjuvant chemotherapy /recurrent cancer without adjuvant chemotherapy and ECOG PS 0/1/2"	
Participants	102 patients with 51 patients each in Slri and SPac arms Median age: 64 (Slri) and 62 (SPac) Advanced/recurrent ratio: 40/11 in both arms "no prior chemotherapy, except adjuvant chemotherapy completed four weeks or more before entry"	
Interventions	Schedule of S-1 was not similar in both arms Arm A: Slri: Irinotecan (i.v.) over 1.5 h at 80 mg/m ² on day 1 and 15, while 40 mg/m ² S-1 (Taiho Pharmaceutical, Tokyo, Japan) was orally administered twice daily for three weeks from days 1-21 followed by a two-week pause. Arm B: SPac: Paclitaxel was administered i.v. over 1 h at a dose of 50 mg/m ² on day 1 and 8, while 40 mg/m ² S-1 was orally administered at twice-daily for two weeks from day1-14 followed by a one-week pause.	
Outcomes	Primary endpoint: Overall response rate Secondary endpoints: Progression-free survival, overall survival and safety	

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Unclear risk	Not stated
Incomplete outcome data (attrition bias)	Low risk	(All outcomes analysed for all patients)

Sugimoto 2014 (Continued)
 efficacy

Incomplete outcome data (attrition bias) safety	Low risk	(All outcomes analysed for all patients)
Selective reporting (reporting bias)	Low risk	Expected endpoints included in reporting
Other bias	High risk	High risk Schedule of S-1 not similar in two arms
Blinded review of CT/MRI-scans?	Low risk	("All radiological assessments were confirmed by extratumoral review")

Thuss-Patience 2005

Methods	Multicentre RCT 2 arms Quality score: A
Participants	n = 90 Median age: 62.5 years Metastatic disease: 98% ECOG 2-3: 3%
Interventions	DF: docetaxel 75 mg/m ² d 1 + FU 200 mg/m ² /day as a 24-hour continuous infusion d 1-21, repeated at d 22 versus ECF: epirubicin 50 mg/m ² d 1 + cisplatin 60 mg/m ² d 1 + FU 200 mg/m ² /as a 24-hour continuous infusion d 1-21, repeated at d 22
Outcomes	Tumour response Median survival Median time to progression Toxicity
Notes	Phase II study The trial was not intended and not statistically powered to perform a head-to-head comparison of response rate and toxicity of the 2 treatment arms. ECF serves as an internal control arm to avoid selection bias.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Random numbers
Allocation concealment (selection bias)	Low risk	Central allocation
Incomplete outcome data (attrition bias) efficacy	Low risk	Analysis of the full analysis set

Thuss-Patience 2005 (Continued)

Incomplete outcome data (attrition bias) safety	Low risk	Analysis of the full analysis set
Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	High risk	The trial was not intended and not statistically powered to perform a head-to-head comparison of response rate and toxicity of the 2 treatment arms, ECF serves as an internal control arm to avoid selection bias.
Blinded review of CT/MRI-scans?	Low risk	Tumour response was assessed together with an independent radiologist

Van Cutsem 2006

Methods	Multicentre RCT 2 arms Quality score: D
Participants	n = 445 Median age: 55 years Metastatic disease: 96.5%
Interventions	DCF: docetaxel 75 mg/m ² + cisplatin 75 mg/m ² d 1 + 5-FU 750 mg/m ² /d as a 24-hour continuous infusion d 1-5, repeated at d 22 versus CF: cisplatin 100 mg/m ² d 1 + 5-FU 1.000 mg/m ² /d as a 24-hour continuous infusion on d 1-5, repeated at d 29
Outcomes	Median overall survival and time to progression Tumour response Toxicity Quality of life
Notes	—

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Low risk	Central allocation
Incomplete outcome data (attrition bias) efficacy	Low risk	Analysis of the full-analysis set
Incomplete outcome data (attrition bias) safety	Low risk	Analysis of the full-analysis set

Van Cutsem 2006 (Continued)

Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	Unclear risk	N/A
Blinded review of CT/MRI-scans?	Low risk	All radiologic assessments were reviewed by an external response review committee and were assessed by WHO criteria

Van Cutsem 2015

Methods	Prospective, multinational, randomised, phase II study Fifty-two sites in the USA and 11 countries in Europe screened and randomised patients	
Participants	The majority (69%) of patients were male; mean age was 59 years	
Interventions	3-arm study: docetaxel/oxaliplatin (TE), docetaxel/oxaliplatin/5-FU (TEF), and docetaxel/oxaliplatin/capecitabine (TEX)	
Outcomes	PFS OS ORR Safety	
Notes	"Tumour response was evaluated every 8 weeks and classified according to best overall response (World Health Organization criteria). Responses were confirmed by two evaluations conducted ≥ 4 weeks apart"	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not clear what method
Allocation concealment (selection bias)	Low risk	RandomiSed centrally using an interactive voice response system
Incomplete outcome data (attrition bias) efficacy	Low risk	"primary efficacy population was the full analysis population (FAP: all randomized and treated patients analysed in the arm to which they were randomized), with supportive analyses conducted using the intent-to-treat (ITT: all randomized patients) and per protocol (PP: assessable patients [received study treatment and had ≥ 1 post-baseline tumour assessment] without any major protocol violation) populations"
Incomplete outcome data (attrition bias) safety	Low risk	"primary efficacy population was the full analysis population (FAP: all randomized and treated patients analysed in the arm to which they were randomized), with supportive analyses conducted using the intent-to-treat (ITT: all randomized patients) and per protocol (PP: assessable patients [received study treatment and had ≥ 1 post-baseline tumour assessment] without any major protocol violation) populations"

Van Cutsem 2015 (Continued)

Selective reporting (reporting bias)	Low risk	PFS, OS, ORR, safety
Other bias	Unclear risk	N/A
Blinded review of CT/MRI-scans?	Unclear risk	Not stated

Wang 2013

Methods	Single-centre RCT 2 arms Quality score: B
Participants	n = 82 participants with metastatic or locally recurrent gastric cancer ECOG 2: 8.5%
Interventions	S-1 +paclitaxel: S-1 depending on body surface area (BSA < 1.25 m ² : 80 mg/d; BSA 1.25 to < 1.5 m ² : 100 mg/d; BSA > 1.5 m ² , 120 mg/d twice daily) d 1-14, paclitaxel 60mg/m ² d 1,8,15, repeated at d 29 versus S-1: S-1 depending on body surface area (BSA < 1.25 m ² : 80 mg/d; BSA 1.25 to <1.5 m ² : 100 mg/d; BSA > 1.5 m ² , 120 mg/d twice daily) d 1-14, repeated at d 29
Outcomes	Overall survival Progression-free survival Response rate Toxicity
Notes	Second-line therapy with cisplatin or irinotecan in more than half of the patients

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Computer-generated, random sequence
Allocation concealment (selection bias)	Low risk	Consecutively numbered opaque sealed envelopes
Incomplete outcome data (attrition bias) efficacy	Low risk	No patients lost to follow-up and all outcomes could be confirmed
Incomplete outcome data (attrition bias) safety	Low risk	No patients lost to follow up and all outcomes could be confirmed
Selective reporting (reporting bias)	Low risk	OS, PFS, ORR, safety

Wang 2013 (Continued)

Other bias	Unclear risk	N/A
Blinded review of CT/MRI-scans?	High risk	Tumor assessment was undertaken with CT or MRI consistently every 2 months by principal investigators

Wang 2016

Methods	Multicentre, prospective, randomised, open-label, phase III trial
Participants	Histo: 15% Signet ring, 3% Others
Interventions	"Untreated advanced gastric cancer patients randomly received docetaxel and cisplatin at 60 mg/m ² (day 1) followed by fluorouracil at 600 mg/m ² /day (days 1–5; mDCF regimen) or cisplatin at 75 mg/m ² (day 1) followed by fluorouracil at 600 mg/m ² /day (days 1–5; CF) every 3 weeks"
Outcomes	"The primary end point was progression-free survival (PFS). The secondary end points were OS, overall response rate (ORR), time-to-treatment failure (TTF), and safety"
Notes	NCT00811447

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated what method was used. "Random assignment was centralized and stratified for center, liver metastases, prior gastrectomy, Karnofsky performance status (KPS) (80 or above vs below 80), and weight loss during the previous 3 months (5 % or less vs more than 5 %)"
Allocation concealment (selection bias)	Low risk	Random assignment was centralised
Incomplete outcome data (attrition bias) efficacy	Low risk	All patients who received treatment were included in full analysis set
Incomplete outcome data (attrition bias) safety	Low risk	All patients who received treatment were included in full analysis set
Selective reporting (reporting bias)	Low risk	Endpoints were all reported
Other bias	Unclear risk	N/A
Blinded review of CT/MRI-scans?	High risk	"Tumor response and PFS were evaluated by investigators, and although much effort has been done to limit the bias in the evaluation of these parameters..."

Webb 1997

Methods	Multicentre RCT 2 arms
---------	---------------------------

Webb 1997 (Continued)

Quality score: B

Participants	n = 199 participants with adenocarcinoma of the stomach or gastro-oesophageal junction Metastatic disease: 85% ECOG 2-3: 26%
Interventions	ECF: epirubicin 50 mg/m ² d 1, repeated at d 22; cisplatin 60 mg/m ² d 1, repeated at d 22, 5-FU CI 200 mg/m ² d 1-21, repeated at d 22 versus FAMTX: methotrexate 1500 mg/m ² d 1; 5-FU 1500 mg/m ² d 1; Lv 30 mg p.o. every 6 hours d 2, 3; adriamycin 30 mg/m ² d 15, repeated at d 28
Outcomes	Median survival Response rates Toxicity Quality of life
Notes	The entire study included 274 participants with inoperable adenocarcinoma or undifferentiated carcinoma of the oesophagus, oesophagogastric junction or stomach. Results for participants with adenocarcinoma of the stomach or gastro-oesophageal junction only were provided by the corresponding author and included in the meta-analysis.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Query to first author was not answered
Allocation concealment (selection bias)	Unclear risk	Query to first author was not answered
Incomplete outcome data (attrition bias) efficacy	Low risk	ITT per correspondence
Incomplete outcome data (attrition bias) safety	Low risk	ITT per correspondence
Selective reporting (reporting bias)	Low risk	Report includes all expected outcomes
Other bias	Unclear risk	N/A
Blinded review of CT/MRI-scans?	High risk	Not done according to correspondence

Wu 2015

Methods	Pilot randomised-controlled study
Participants	"158 participants were initially screened, of whom 86 were excluded. Of these 86 patients, 73 did not fulfill the study criteria and 13 declined to participate. The remaining 72 patients (36 treated with SC and 36 treated with C) were entered into the study"

Wu 2015 (Continued)

"The mean age of the patients was 64.1 years in the SC group and 62.7 years in the C group. The performance status was 0 for 41.7% of patients treated with SC and 44.4% of patients treated with C and it was 1 for 58.3% of patients treated with SC and 44.4% of patients treated with C. The primary lesion was 55.6% in the SC group and 50.0% in the C group. Histological types were intestinal (58.3% in the SC group and 61.1% in the C group), diffuse (36.1% in the SC group and 30.6% in the C group), and others (5.6% in the SC group and 8.3% in the C group). The diagnosis was AGC (86.1% in the SC group and 83.3% in the C group) and relapse gastric cancer (13.9% in the SC group and 16.7% in the C group)."

Interventions	"Patients in the C group received cisplatin 75 mg/m ² intravenously over 1–3 h on day 1 and then at 4-week intervals. In addition to receiving the same intervention as the C group, patients in the SC group were also administered S-1 on days 1–14 of a 21-day cycle. The daily dose of S-1 was assigned according to the body surface area as follows: less than 1.25 m ² , 40 mg two times daily; more than or equal to 1.25 m ² and less than 1.5 m ² , 50 mg two times daily; and more than or equal to 1.5 m ² , 60 mg two times daily."
Outcomes	OS PFS Adverse events

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	"Patients were stratified using the block randomization method of the SAS package (version 8.2; SAS Institute Inc., Cary, North Carolina, USA) by a statistician with no clinical involvement in this study"
Allocation concealment (selection bias)	Low risk	"The allocation was concealed in sequentially numbered, opaque, sealed envelopes containing the randomization assignments"
Incomplete outcome data (attrition bias) efficacy	Low risk	All participants could undergo evaluations for efficacy and safety.
Incomplete outcome data (attrition bias) safety	Low risk	All participants could undergo evaluations for efficacy and safety.
Selective reporting (reporting bias)	High risk	Overall response rates were compared although it did not appear to be pre-specified. Furthermore, details of how response was assessed were not provided.
Other bias	High risk	<p>Study did not clearly specify minimum required sample size - "Sample size was calculated on the basis of an expected 15% difference between the two groups, with 80% power and a two-sided [alpha] value of 0.05"</p> <p>Comments from statistician BCT: The paper is not clear about how the sample size is being estimated. In particular, we do not have information on the HR or the survival probability of the control group. I assume the 15% difference refers to absolute difference. I try to work out this difference from the KM curve of Fig 2. If we assume a 1 year estimate, then the survival probability is approx. 25% vs 40% (HR approx. 0.66). If we assume a 0.5 year estimate, then the survival probability is 75% vs 90% (HR approx. 0.37). This corresponds to a total sample size of about 300 and 200 respectively, assuming a two-sided test at 5% level and a power of 80%. If we estimate the HR from the median OS that</p>

Wu 2015 (Continued)

is reported, the HR is approx. 0.8, and so we would expect an even larger sample size.

Blinded review of CT/MRI-scans?	Unclear risk	Not stated
---------------------------------	--------------	------------

Yamada 2015

Methods	Randomised, open-label, multicentre phase III study
Participants	685 patients were enrolled; 343 and 342 patients were randomly assigned to SOX or CS
Interventions	"In CS, S-1 was given orally twice daily for the first 3 weeks of a 5-week cycle. The dose was 80 mg/day for body surface area (BSA) <1.25 m ² , 100 mg/day for BSA ≥1.25 to <1.5 m ² , and 120 mg/day for BSA ≥1.5 m ² . Cisplatin was administered at 60 mg/m ² as an i.v. infusion with adequate hydration on day 8 of each cycle [9]. In SOX, S-1 was given as the same way for the first 2 weeks of a 3-week cycle. Oxaliplatin at 100 mg/m ² was infused for 2 h i.v. on day 1 of each cycle"
Outcomes	The primary end points were noninferiority in progression-free survival (PFS) and relative efficacy in overall survival (OS) for SOX using adjusted hazard ratios (HRs)

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Minimisation method.
Allocation concealment (selection bias)	Low risk	The randomisation sequence was generated by an independent team from the trial sponsor and investigators
Incomplete outcome data (attrition bias) efficacy	Low risk	ITT
Incomplete outcome data (attrition bias) safety	Low risk	ITT
Selective reporting (reporting bias)	Low risk	PFS, OS, tumor responses, Adverse events
Other bias	Unclear risk	"During the study period, we did not test HER2 expression in tumors and could not know its exact influence on our results. The proportion of patients who received trastuzumab after the study treatment was small (<10%) and similar in both groups. Therefore, trastuzumab treatment would not seem to impact on comparing OS between both groups." "Patients who were alive and free of progression (i.e. second-line treatment was started due to any cause) were regarded as censored cases at the date of the last assessment"

Yamada 2015 (Continued)

"In February 2011, it appeared to be difficult to achieve the required number of events within the preplanned timetable, and the target number of patients was revised to 680 according to the predefined procedure in the protocol"

Blinded review of CT/MRI-scans?	Low risk	"All images for PFS and tumor responses were reviewed by an independent review committee, according to the Response Evaluation Criteria In Solid Tumors (RECIST) version 1.0"
---------------------------------	----------	---

Yamamura 1998

Methods	Multicentre RCT 2 arms Quality score: C
Participants	n = 71
Interventions	MTX/5-FU/THP: methotrexate 50 mg/m ² , 5-FU 650 mg/m ² , pirarubicin 20 mg/m ² d 1, repeated at d 15 versus 5-FU: 5-FU 650 mg/m ² d 1, repeated at d 15
Outcomes	Median survival Toxicity
Notes	Study translated from Japanese

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not stated
Allocation concealment (selection bias)	Low risk	Centralised. Phone call
Incomplete outcome data (attrition bias) efficacy	Low risk	71/74 evaluable
Incomplete outcome data (attrition bias) safety	Low risk	71/74 evaluable
Selective reporting (reporting bias)	High risk	No response rates available JCOG criteria for toxicity
Other bias	Unclear risk	Study translated from Japanese
Blinded review of CT/MRI-scans?	Unclear risk	N/A

Abbreviations:

BSA: body surface area
 BSC: best supportive care
 CI: continuous infusion

CT: computed tomography
 d: day
 dFUR: 5-Deoxyfluoridine
 ECOG: Eastern Cooperative Oncology Group
 5-FU: 5-Fluorouracil
 FA: folinic acid
 h: hour
 IQR: interquartile range
 ITT: intention-to-treat
 i.v.: intravenous
 KI: Karnofsky-Index
 KPS: Karnofsky performance status
 Lv: leucovorin
 NCI-CTC: National Cancer Institute Common Toxicity Criteria
 OS: overall survival
 p.o.: per os (orally)
 PFS: progression-free survival
 QoL: quality of life
 RCT: randomised controlled trial
 RR: response rate
 TTP: time to progression
 UFT: uracil/ftorafur
 w: week
 WHO: World Health Organization
 If not stated differently, all drugs were given as intravenous bolus or short infusion (duration max. 2 hours)

Characteristics of excluded studies [ordered by study ID]

Study	Reason for exclusion
Ahn 2002	According to information provided by the first author about 2/3 of included participants had undergone gastric resection and were treated in adjuvant/additive intention.
Ajani 2002	Article about study published elsewhere (Van Cutsem 2006).
Ajani 2006	Non-randomised phase II study.
Akagi 2010	Other indication (patients underwent macroscopically curative resection).
Akazawa 1985	Study not randomised.
Andrić 2012	Interventions not relevant for any of the comparisons specified.
Anonymous 1979	Not all participants were randomised (some were directly assigned to 1 treatment arm according to their prior chemotherapy). Systematic cross-over between treatment arms.
Anonymous 1982	Study began as three-arm comparison of FAME, FAMi and FIME regimens. FMe was added to randomization later. ICRF-159 =dextrazoxaneL None of the possible permutations between these regimens fit our study comparisons
Anonymous 1983	18 of 82 included participants were crossed over (< 10 %)
Anonymous 1984	Compared FA, FAME, and FAMi. None of the possible permutations between these regimens fit our study comparisons.
Aoyama 1981	Study not randomised

Study	Reason for exclusion
Bajetta 1998	Participants not randomised to different chemotherapy regimens, but to one regimen (FEP) with or without GM-CSF
Baker 1976	Participants with different advanced gastrointestinal cancers were included in the study. No separate results given for gastric cancer participants.
Balana 1990	Preliminary publication.
Berenberg 1989	Final publication: Berenberg 1995 (excluded).
Berenberg 1995	Compares different single-agent chemotherapy regimens.
Beretta 1983	Published as abstract only. No further information obtained by contacting the first author.
Beretta 1989	Published as abstract only. No further information obtained by contacting the first author.
Berglund 2006	Other indications.
Bi 2011	Cross-over study.
Bjerkset 1986	Study included participants with different advanced gastrointestinal cancers. No separate results given for gastric cancer participants.
Bruckner 1986	Preliminary publication.
Bugarolas 1975	Study published as abstract only with insufficient data available. According to information provided by the first author no further analysis was performed as the study was terminated early due to slow accrual.
Bugat 2003	Secondary publication.
Buroker 1979	Does not compare different intravenous combination chemotherapy regimens but 1 intravenous regimen with or without an additional oral chemotherapy (methyl-CCNU).
Cai 2011	Not an RCT, retrospective analysis of an RCT.
Cascinu 1994	Final publication: Cascinu 1995 (excluded).
Cascinu 1995	Participants not randomised to different chemotherapy regimens but to reduced glutathione or placebo to prevent cisplatin-induced neurotoxicity.
Cascinu 1996	Participants not randomised to different chemotherapy regimens but to different doses of G-CSF as supportive therapy.
Chau 2013	Testing VEGF inhibitor (ramucirumab) as second-line therapy.
Chen 2011	Gimeracil + oteracil, neither contains a 5-FU prodrug nor is equivalent to S-1. So this combination does not fit any comparison.
Chlebowski 1979	One study arm was treated with oral 5-FU.
Chlebowski 1985	Insufficient information for calculation of HRs given in the publication or provided by the author.
Chu 2006	Missing information to calculate HRs for OS

Study	Reason for exclusion
Chung 2011	Not an RCT.
Coates 1984	Included 108 participants with advanced cancer (only 5 gastric cancer patients).
Cocconi 1982	Systematic cross-over between study groups (< 10 % of included participants).
Cocconi 1992	Final publication: Cocconi 1994 (included).
Colucci 1991	Final publication: Colucci 1995 (included).
Constenla 2002	Study not randomised.
Coombes 1994	Compares different single agents.
Cullinan 1993	Final publication: Cullinan 1994 (included).
Cunningham 2008	Wider indication (inclusion of patients with squamous cell carcinoma of the oesophagus as well).
De Lisi 1985	Final publication: De Lisi 1986 (included).
De Lisi 1988	Second publication to De Lisi 1986 (included).
Diaz-Rubio 1991	Preliminary publication.
Douglass 1983	Final publication: Douglass 1984 (excluded).
Douglass 1984	Not all participants were randomised. Some were directly assigned to 1 study arm according to their prior treatment.
Duffour 2006	Interventions not relevant for any of the comparisons specified.
Figoli 1991	Study not randomised. Two consecutive regimens were compared.
Ford 2014	Second-line CTX.
Fuchs 2014	Testing VEGF inhibitor (ramucirumab) as second-line therapy.
Fujii 1983	Study not randomised.
Furue 1985	Schizophyllan: mushroom polysaccharide with immunomodulatory properties (biologic therapy), no chemotherapy.
Furukawa 1995	Although described as advanced gastric cancer in the title, according to the text of the article the study compared different adjuvant chemotherapy regimens.
Gao 2010	inclusion pf patients with squamous epithelium carcinoma.
Gioffre 1984	Study not randomised.
Glimelius 1994	Final publication: Glimelius 1997 (excluded).
Glimelius 1995	Study includes participants with different inoperable cancers. Cross-over from BSC to chemotherapy arm permitted. Primary aim: cost-effectiveness.
Glimelius 1997	Study compares chemotherapy versus best supportive care. Provision of chemotherapy upon request in the best supportive care group was requested by the research ethics committee, and 12 of

Study	Reason for exclusion
	30 participants (19.6%) randomised to best supportive care finally received chemotherapy (cross-over).
Goseki 1995	Participants were not randomised between different chemotherapy regimens but to methionine-depleted total parenteral nutrition versus a conventional amino acid solution.
Grau 1988	All participants were treated with chemotherapy after resection (no advanced/metastatic disease).
Grieco 1984	Study not randomised.
Gubanski 2010	Prescheduled cross-over between study arms after four courses.
Guimbaud 2014	Considerable use of second-line therapy.
Gupta 1982	Insufficient data on survival available.
Haas 1983	Cross-over after failure was encouraged.
Hawkins 2003	Preliminary data.
Hoffman 1998	Retrospective analysis of clinical benefit and quality of life in participants with different inoperable gastrointestinal cancers.
Icli 1993	Study not randomised.
Imada 1999	Includes participants treated in adjuvant intention after curative resection of gastric cancer.
Inoue 1989	Participants with ascites were treated with intraperitoneal chemotherapy.
Jeung 2011	Interventions not relevant for any of the comparisons specified.
Kang 2007	Interventions not relevant for any of the comparisons specified.
Kelsen 1990	Preliminary publication.
Kilickap 2011	Not an RCT.
Kim 1991	Final publication: Kim 1993 (excluded).
Kim 1993	According to information provided by author (YSP) allocation was done by alternation (not truly randomised).
Kim 2012	Interventions not relevant for any of the comparisons specified.
Kim 2013	Meta-analysis is examining second-line therapy vs BSC.
Kitamura 1995	Study compares the effect of different amino-acid solutions (methionine-free amino-acid solution versus commercial amino-acid solution) in addition to 5-FU chemotherapy.
Koizumi 1996	Compares different dosages of cisplatin (60 mg and 80 mg) within the same chemotherapy regimen.
Koizumi 2004	Interventions not relevant for any of the comparisons specified.
Koizumi 2012	Not an RCT.

Study	Reason for exclusion
Koizumi 2013	Testing irinotecan plus cisplatin versus irinotecan as second-line therapy.
Kolaric 1986	Participants with stomach and rectosigmoid cancer included. No separate information about gastric cancer provided.
Kondo 2000	Study uses oral 5-FU (no intravenous chemotherapy, varying bioavailability).
Kono 2002	Study treatment: adoptive immunotherapy.
Kornek 2002	Does not compare different chemotherapy regimens but the same chemotherapy with or without G-CSF and/or erythropoietin.
Kosaka 1995	One group was treated with intra-arterial chemotherapy.
Kovach 1974	Insufficient data for calculation of HRs given (P value missing).
Kuitunen 1991	Final publication: Pyrhönen 1995 (included).
Kurihara 1991	<p>Compared: tegafur + mitomycin C (FTM), uracil-tegafur + mitomycin C (UFTM), 5'-deoxy-fluorouridine + cisplatin (5'P), etoposide + doxorubicin + cisplatin (EAP), and 5-fluorouracil + cisplatin (FP). Firstly,</p> <p>none of the possible permutations between these regimens fit our study comparisons. Secondly, the Ohkuwa 2000 paper resembles a "<i>pooled</i>" analysis (of other RCTs such as Kurihara 1991) rather than an <i>original</i> RCT.</p>
Kurihara 1995	Participants randomised to either methionine-free or commercial amino-acid solution, with the same (mitomycin C/fluorouracil) chemotherapy in both groups.
Kurihara 1995a	Final publication: Koizumi 1996 (excluded).
Lacave 1985	Final publication: Lacave 1987 (excluded).
Lacave 1987	Does not compare different intravenous chemotherapy regimens but 1 intravenous chemotherapy with or without an additional oral chemotherapy (methyl-CCNU).
Lee 2008	Interventions not relevant for any of the comparisons specified.
Lee 2012	Gastrointestinal endoscopy.
Levard 1998	Participants had oesophageal squamous cell carcinoma.
Li 2002	Drug under investigation was never approved, and the regimen does not fit any of our 10 comparisons. Also their trial was not solely done in gastric cancer patients, but also included patients with colorectal, oesophageal, and liver cancer.
Li 2007	Missing information to calculate HRs for OS.
Li 2011	Not relevant for any of the comparisons.
Li 2013	Testing second-line apatinib.
Lim 2011	Not relevant for any of the comparisons.
Livstone 1977	Radiochemotherapy is compared to systemic intravenous chemotherapy.

Study	Reason for exclusion
Lordick 2013	Testing whether cetuximab (EGFR targeted mAB) should be included as first-line in combination with capecitabine and cisplatin.
Lorenzen 2007	Not a randomised trial.
Luelmo 2006	Interventions not relevant for any of the comparisons specified.
Malik 1990	Study not randomised. Includes participants with gastric and colorectal cancer.
Maruta 2007	Second-line chemotherapy.
Massuti 1994	Published as abstract only. No further information obtained by contact with the first author.
Massuti 1995	Published as abstract only. No further information obtained by contact with the first author.
Mochiki 2012	Interventions not relevant for any of the comparisons specified.
Moertel 1976	Insufficient information for calculation of HRs given.
Moertel 1979	The combination therapy arm consisted of only 1ne oral agent (methyl-CCNU, no intravenous chemotherapy), in addition to 5-FU which was used as single-agent.
Moertel 1979a	Not all participants were randomised (some were directly assigned to 1 treatment arm according to their prior treatment). Systematic cross-over between study groups.
Moore 2005	Secondary publication.
Mustacchi 1997	Study included participants with different advanced cancers (only 3 patients with gastric cancer).
Nakajima 1984	Study compares different adjuvant chemotherapies.
Nakao 1983	Study treatment included schizophyllan (mushroom polysaccharide with immunomodulatory effects, biological therapy).
Nakashima 2008	The treatment for each patient was decided by the patient's choice or randomisation (not a randomised trial).
Niitani 1987	Compares different modes of application of the same chemotherapy (continuous and intermittent oral administration of 5´ deoxy-5-fluorouridine), not different chemotherapy regimens.
Nordin 2001	Study presents different interpretations of quality of life data from a previously performed study (Glimelius 1997). The study included advanced gastric cancer participants, but was excluded because of cross-over.
Novik 1999	Compares different single agents.
Ohtsu 2011	Testing whether bevacizumab (VEGF targeted mAB) should be included in first-line chemotherapy combinations.
Okines 2010	Not an RCT.
Osawa 1996	Study treatment: adjuvant chemo-immunotherapy.
Pannettiere 1984	Compares 2 modes of application (sequential versus simultaneous) of 1 chemotherapy regimen (FAM).

Study	Reason for exclusion
Park 2004	Interim analysis of Park 2006 .
Park 2006	Interventions not relevant for any of the comparisons specified.
Park 2008	Interventions not eligible for any of the comparisons specified.
Popliela 1982	Study treatment: chemo-immunotherapy.
Popov 1999	Final publication: Popov 2000 (excluded).
Popov 2000	Compares different applications of doxorubicin (bolus versus 8-hour infusion) in the same chemotherapy regimen (EAP), not different chemotherapy regimens.
Pozzo 2004	Interventions not relevant for any of the comparisons specified.
Pyrhonen 1992	Final publication: Pyrhönen 1995 (included).
Queisser 1984	Compared 5-fluorouracil and carmustine with or without adriamycin. This does not fit our study comparisons.
Rake 1979	39 of 77 included participants receiving chemotherapy as additive therapy after non-curative resection with histological evidence of residual disease.
Roth 1994	Final publication: Roth 1999 (included).
Roth 1995	Final publication: Roth 1999 (included).
Roth 1997	Final publication: Roth 1999 (included).
Sakata 1982	Participants were treated with OK-432 (streptococcal preparation, biological response modifier).
Sakata 1988	Study included participants with various gastrointestinal tumors, which were treated with a biological therapy.
Sakata 1992	Study included participants with different adenocarcinomas.
Sasagawa 1994	Insufficient information for calculation of HRs given.
Sasaki 1989	Publication presents only preliminary results of this study. No information about final results provided by the first author.
Sasaki 1990	Study not randomised.
Sasaki 1992	Study includes participants with colorectal and gastric cancer. Insufficient information given about results in gastric cancer.
Sasaki 1995	Study includes participants with colorectal and gastric cancer. Insufficient information given about results in gastric cancer.
Sato 1991	Study summarises the experience of angiotensin II-induced hypertension to enhance drug delivery for chemotherapy. Participants were not randomised to different chemotherapy regimens.
Sato 1995	Participants were not randomised to different chemotherapy regimens but to angiotensin II-induced hypertension to enhance chemotherapy effects or control.

Study	Reason for exclusion
Satoh 2013	Text needed.
Satoh 2014	Testing whether lapatinib (HER2 inhibitor) should be used in second-line therapy in HER2-amplified Asian population.
Schmid 2003	Study not randomised.
Shen 2009	No information on overall survival.
Shin 2007	Interventions not relevant for any of the comparisons specified.
Shinoda 1995	Reporting standards to be insufficient and/or the dose schedule to be different from other studies.
Shu 1999	Study compared the effect of intraperitoneal combined with intravenous to only intravenous chemotherapy.
Shudong 1996	Insufficient information for calculation of HRs given.
Smith 1983	Study included various advanced adenocarcinomas of the gastrointestinal tract (only 5 participants with gastric cancer).
Sun 2004	Interventions not relevant for any of the comparisons specified (radiochemotherapy).
Sym 2013	A study in participants refractory to first-line chemotherapy - thus no first-line treatment.
Taal 1990	Study not randomised.
Taguchi 1985	Study treatment includes lentinan (mushroom polysaccharide with immunomodulatory properties, biological therapy).
Takahashi 1991	Study participants had resectable tumours. One group received intraarterial chemotherapy.
Tebbutt 2002	Not eligible for any of the comparisons specified.
Tebbutt 2007	Not relevant for any of the comparisons (docetaxel in both treatment arms).
Tebbutt 2010	Included esophageal cancer patients.
Thuss-Patience 2011	Second-line therapy.
Tsushima 1991	Study treatment includes OK (streptococcal preparation, biologic response modifier).
Van Cutsem 2009	Targeted therapy.
Vanhoefler 2000	Compared methotrexate, fluorouracil, and doxorubicin versus etoposide, leucovorin, and fluorouracil versus infusional fluorouracil and cisplatin. None of the possible permutations between these regimens fit our study comparisons
Vaughn 1980	Study included participants with advanced gastrointestinal malignancies (only 11 participants with gastric cancer).
Vestlev 1990	Inconclusive data (time to progression and survival identical).
Villar 1987	Study included participants (13 of 46) with only microscopic disease in the resection margins.

Study	Reason for exclusion
Voznyi 1978	One study arm consisted only of oral chemotherapy (CCNU). Two other of 5 study arms differed from others only in the addition of oral CCNU (does not compare different intravenous chemotherapy regimens).
Wadler 2002	Study treatment includes interferon and filgrastim (biological therapy).
Wakui 1983	Participants had various gastrointestinal malignancies. Therapy includes levamisole (antihelmintic drug with immunomodulatory properties).
Wakui 1983a	Participants had various gastrointestinal malignancies. Therapy included levamisole (antihelmintic drug with immunomodulatory properties).
Wakui 1986	Study treatment includes lentinan (mushroom polysaccharide with immunomodulatory properties, biological therapy).
Wang 2007	Not relevant for any of the comparisons.
Waters 1999	Paper reports long-term follow-up of the study published by Webb 1997 (included)
Wilke 2014	Testing VEGF inhibitor (ramucirumab) as second-line therapy.
Wils 1991	Compared 5-fluorouracil, adriamycin and methotrexate vs 5-fluorouracil, adriamycin and mitomycin-C. This does not fit our study comparisons
Wils 1994	Published as abstract only, no further information about final results obtained by contacting the first author.
Xu 2013	Tested recombinant human endostatin, and regimen does not fit our comparisons.
Yamada 1994	According to information provided by the first author the study was not randomised.
Yin 1996	Study treatment includes Elemene (product isolated from the Chinese medical herb <i>Rhizoma zedoariae</i> , biological therapy).
Yoshida 2003	Compares the feasibility of personalised chemotherapy (according to the expression of molecular markers) with standard therapy.
Yoshikawa 2011	Tested a chemoimmunotherapy combination.
Yoshino 2007	Intervention not eligible for any of the comparisons specified.
Yun 2010	Intervention not eligible for any of the comparisons specified.
Zhao 2009	missing information to calculate HRs for OS.
Zironi 1992	Final publication: Cocconi 1994 (included)

5-FU: 5-Fluorouracil

A: adriamycin (also known as doxorubicin)

BSC: best supportive care

CTX: chemotherapy

FA: folinic acid

FAMe: 5-FU 325mg/m² d 1-5, A 40mg/m² d 1 until d 36, Me 110mg/m² orally at d 1 repeated at d 71

FAMi: 5-FU 275mg/m² d 1-5, A 30mg/m² d 1 until d 36, M 10mg/m² d1 repeated at d 71

FIMe: 5-FU 325mg/m² d 1-5 repeated at d 36, ICRF-159 500mg/m² orally at d 2-4 and d 36-38, Me 110mg/m² orally at d 1 repeated at d 71

FMe: 5-FU 300 mg/m² d 1-5 repeated at d 36, Me 175 mg/m² orally at d 1 repeated at d 50

Chemotherapy for advanced gastric cancer (Review)

Copyright © 2017 The Cochrane Collaboration. Published by John Wiley & Sons, Ltd.

G-CSF: granulocyte-macrophage colony-stimulating factor

HR: hazard ratio

ICRF: razoxane

M: mitomycin C

Me: semustine (also known as methyl-CCNU)

OS: overall survival

RCT: randomised controlled trial

VEGF: vascular endothelial growth factor

Characteristics of ongoing studies [ordered by study ID]

Elsaid 2005

Trial name or title	Randomized phase III trial of docetaxel, carboplatin and 5FU versus epirubicin, cisplatin and 5FU for locally advanced gastric cancer (final publication not found)
Methods	RCT 2 arms
Participants	N = 64
Interventions	Arm A: docetaxel (75 mg/m ² d 1) + carboplatin (AUC6 d2) + continuous infusion 5FU (1200 mg/m ² /day d1-3), repeated at d 21 versus Arm B: Epirubicin (50 mg/m ² d 1) + cisplatin (60 mg/m ² d1)+ continuous infusion 5FU (200 mg/m ² /day d 1-21) Prophylactically G-CSF day 4-9 to all participants
Outcomes	Toxicity, tumour response, progression-free and overall survival
Starting date	1999
Contact information	Amr Abdelaziz Elsaid amrelsaid@yahoo.com Alexandria University, Alexandria, Egypt
Notes	This study is currently published as abstract only. Participants were enrolled between 1999 and 2004.

Higuchi 2012

Trial name or title	Randomized phase III study of S-1 plus oxaliplatin versus S-1 plus cisplatin for first-line treatment of advanced gastric cancer
Methods	RCT 2 arms
Participants	N = 685 with unresectable advanced or recurrent gastric cancer, PS 0-2, age ≥ 20 years
Interventions	Arm A (SOX): (oral S-1 40 mg/m ² twice a day d 1-14 + oxaliplatin 100 mg/m ² iv day 1, repeated at 3 weeks) versus

Higuchi 2012 (Continued)

Arm B (SP): (oral S-1 40 mg/m² twice a day d 1- 21 + cisplatin 60 mg/m² iv day 8, repeated at 5 weeks)

Outcomes	Progression-free survival, response rate (RR), safety, and length of hospital stay per cycle
Starting date	
Contact information	Yakult Honsha Co., Ltd., Clinical Development, Medical Development Department, Pharmaceutical Division, Tel 813-5550-8966 Fax 813-3248-5502
Notes	The study is registered (JapicCTI-101021) and has recruited 685 participants from January 2010 and October 2011.

Kurihara

Trial name or title	Isovorin: Phase III study
Methods	RCT 2 arms
Participants	Participants with histologically confirmed, advanced gastric adenocarcinoma without prior chemotherapy, PS 0-2, age 20 to 70 years
Interventions	5-FU+ isovorin (active form of leucovorin) versus S-1 alone
Outcomes	Overall survival, response rates, time to progression, safety, quality of life
Starting date	—
Contact information	Minoru Kurihara, M.D., Showa University, Japan
Notes	Sponsor: Wyeth Pharmaceuticals. Final publication not found.

Maiello 2011

Trial name or title	Epirubicin (E) in combination with cisplatin (CDDP) and capecitabine (C) versus docetaxel (D) combined with 5-fluorouracil (5-FU) by continuous infusion as front-line therapy in participants with advanced gastric cancer (AGC)
Methods	RCT 2 arms
Participants	N = 77 with advanced gastric cancer and measurable disease
Interventions	Arm A (ECX): epirubicin (50 mg/m ² d 1) + cisplatin (60 mg/m ² d 1) + capecitabine (625 mg/m ² twice a day, d 1-21) repeated at d 21 versus Arm B (DF): docetaxel (85 mg/m ² d1)+ 5-FU (750 mg/m ² /day, d1-5) qd 21

Maiello 2011 (Continued)

Outcomes	Response rate, toxicity
Starting date	
Contact information	G.Colucci, Istituto Oncologici Bari, Bari, Italy
Notes	Preliminary results of this study are currently published as abstract only. Final publication not found.

NCT01498289

Trial name or title	A randomized phase II pilot study prospectively evaluating treatment for participants based on ERCC1(excision repair cross-complementing 1) for advanced/metastatic oesophageal, gastric or gastroesophageal junction (GEJ) cancer
Methods	Multicentre RCT 2 arms stratification according to ERCC1
Participants	N = 225 with unresectable, advanced or recurrent gastric cancer, age ≥18 years, HER-2 negative
Interventions	Arm A (FOLFOX): 4-FU (bolus 400 mg/m ² , ci 2400 mg/m ²) + LV (400 mg/m ²) + oxaliplatin (85 mg/m ²) d 1, repeated at d 14. Arm B docetaxel (30 mg/m ² on day 1 + 8 qd 21) + irinotecan (65 mg/m ² on day 1 + 8 qd 21).
Outcomes	Progression-free survival (PFS) between high-ERCC1 and low-ERCC1 participants treated with FOLFOX versus irinotecan hydrochloride plus docetaxel, overall survival, response rate, toxicity
Starting date	February 2012
Contact information	Contact: Kimberly Kaberle: kkaberle@swog.org ; Dana Sparks: dsparks@swog.org Principal Investigator: Syma Iqbal
Notes	Sponsors: Southwest Oncology Group and National Cancer Institute (NCI)

NCT01558947

Trial name or title	Peri-operative chemotherapy with ECX (Epirubicin + Cisplatin + Capecitabine) or XP (Capecitabine + Cisplatin) in the treatment of advanced gastric cancer: a randomized, multicentre, parallel vontrol
Methods	Allocation: Randomised Intervention Model: Parallel assignment Masking: Single-blind (participant) Primary Purpose: Treatment
Participants	Gastric cancer participants, ≥ T2 or N +; or staging II, IIIA, IIIB.
Interventions	Experimental: chemotherapy with ECX

Chemotherapy for advanced gastric cancer (Review)

NCT01558947 (Continued)

Preoperative chemotherapy of ECX for 3 cycles (Epirubicin 50 mg/m² on day 1; capecitabine 1000 mg/m², 2 times /day, 1 to 14 days; cisplatin 60 mg/m² on day 1, need hydration, 21 day/cycle), operation after 2~4 weeks, and postoperative chemotherapy of ECX for 3 cycles 4~6 weeks after surgery.

Experimental: chemotherapy with XP

Preoperative chemotherapy of XP for 3 cycles (capecitabine 1000 mg/m², 2 times / day, 1 to 14 days; cisplatin 60 mg/m² on day 1, need hydration, 21 day/cycle), operation after 2~4 weeks, and postoperative chemotherapy of XP for 3 cycles 4~6 weeks after surgery.

Other Name: XP chemotherapy

Outcomes	Relapse-free survival time/rate [Time Frame: 3 years]
Starting date	January 2011
Contact information	Xiangdong Cheng, MD, Zhejiang Cancer Hospital
Notes	

NCT01967875

Trial name or title	A prospective, multi-center, randomized controlled phase 2 trial of optimising platinum-based chemotherapy based on ERCC1 expression as first-line treatment in participants with locally advanced or metastatic gastric cancer
Methods	Multicentre RCT 3 arms
Participants	N = 180, advanced gastric cancer, 18-65 years, KPS 70, measurable disease
Interventions	Active comparator: ERCC1 high expression-group A: Cisplatin 75 mg/m ² , d 1; Capecitabine 1700 mg/m ² /day to 2000 mg/m ² /day d 1-14 repeated at d 21 for 6 cycles. Capecitabine is to be continued until disease progression or intolerable toxicity. Experimental: ERCC1 high expression-group B: Docetaxel 75 mg/m ² , d 1; Capecitabine 1700 mg/m ² /day to 2000 mg/m ² /day on days 1-14 every 21 days, 6 cycles. Capecitabine is to be continued until disease progression or intolerable toxicity. Active comparator: ERCC1 low expression group: Cisplatin 75 mg/m ² , d 1; Capecitabine 1700 mg/m ² /day to 2000 mg/m ² /day d 1-14 repeated at d 21 for 6 cycles. Capecitabine is continued until disease progression or intolerable toxicity.
Outcomes	Progression-free survival, overall survival, objective response rate, disease control rate, duration of response, safety, quality of life
Starting date	July 2013
Contact information	Yunpeng Liu: cmuliuyunpeng@hotmail.com , Jing Liu: liujing_cmu@hotmail.com
Notes	Sponsor: China Medical University, China

NCT02076594

Trial name or title	A randomized phase III study of low-docetaxel oxaliplatin, capecitabine (low-tox) vs epirubicin, oxaliplatin and capecitabine (Eox) In patients with locally advanced unresectable or metastatic gastric cancer
Methods	<p>Phase 3</p> <p>Allocation: Randomised</p> <p>Intervention Model: Parallel assignment</p> <p>Masking: Open-label</p> <p>Primary Purpose: Treatment</p>
Participants	<p>N = 462</p> <p>Age 18 to 69 years</p> <ul style="list-style-type: none"> · Histologically proven diagnosis of adenocarcinoma of the stomach · HER2 negative tumour or HER2+ tumours not qualifying for herceptin therapy · Locally advanced (non resectable) or metastatic gastric cancer
Interventions	<p>Experimental arm A: docetaxel & oxaliplatin & capecitabine</p> <p>Participants will receive cycles every 3 weeks of docetaxel (35 mg/m², intravenous at days 1 and 8 by 1-hour infusion) and oxaliplatin (80 mg/m², intravenous at day 1 by 2-hour infusion) and capecitabine (750 mg/m², oral tablets of 500 mg and 150 mg, x2 daily for 2 weeks)</p> <p>Experimental arm B: epirubicin & oxaliplatin & capecitabine</p> <p>Participants will receive cycles every 3 weeks of epirubicin (50 mg/m², intravenous on day 1 by 2-hour infusion) and oxaliplatin (130 mg/m², intravenous on day 1 by 2-hour infusion) and capecitabine (625 mg/m², oral tablets of 500 mg and 150 mg, x2 daily for 3 weeks)</p>
Outcomes	<p>Overall survival (OS) [Time frame: Measured as the time from randomisation to the date of death from any cause, assessed up to 18 months of follow-up]</p> <p>Progression-free survival (PFS) [Time frame: Measured as the time from randomisation to the date of local or regional progression, distant metastasis, second primary malignancy or death from any cause, whichever comes first, assessed up to 18 months of follow-up]</p> <p>Objective Response Rate (CR + PR) according to RECIST 1.1 guideline [Time frame: Measured as the time from randomisation, assessed up to 18 months of follow-up]</p> <p>Disease control rate: CR + PR + SD lasting > 12 weeks [Time frame: Measured as the time from randomisation, assessed up to 18 months of follow-up]</p> <p>To assess tolerability of the treatments of participants with locally advanced unresectable or metastatic gastric cancer treated with docetaxel plus oxaliplatin plus capecitabine (Arm A) or with epirubicin plus oxaliplatin plus capecitabine (Arm B)</p>
Starting date	January 2013
Contact information	Contact: Roberto Labianca, MD +39 035 2673691 rlabianca@hpg23.it
Notes	

NCT02114359

Trial name or title	Comparison of efficacy and tolerance between combination therapy and monotherapy as first line chemotherapy in elderly participants with advanced gastric cancer: a multicenter randomized phase 3 study
Methods	<p>Multicentre RCT</p> <p>Phase 3</p> <p>Allocation: Randomised</p> <p>Intervention Model: Parallel assignment</p> <p>Masking: Open-label</p> <p>Primary Purpose: Treatment</p>
Participants	N = 332, ≥ 70 years, Eastern Cooperative Oncology Group 0-2, Measurable or evaluable disease, HER-2 negative
Interventions	<p>Experimental: Platinum/fluoropyrimidine combination chemotherapy</p> <ul style="list-style-type: none"> · Drug: Capecitabine/cisplatin <p>Capecitabine/cisplatin (XP) : cisplatin 50 mg/m² (80% dose of 60 mg²m) iv over 15 min D1, capecitabine 1000 mg/m² (80% dose of 1250 mg/m²) orally twice a day D1-14, repeated at 3 weeks</p> <ul style="list-style-type: none"> · Drug: S-1/cisplatin <p>S-1/cisplatin (SP) : cisplatin 50 mg/m² (80% dose of 60 mg²m) iv ov 15min D1, S-1 30 mg/m² (80% dose of 40 mg/m²) orally twice a day D1-14, repeated at 3 weeks</p> <ul style="list-style-type: none"> · Drug: Capecitabine/oxaliplatin <p>Capecitabine+oxaliplatin (XELOX): oxaliplatin 100 mg/m² (80% dose of 130 mg/m²) iv ov 120 min D1, capecitabine 800 mg/m² (80% dose of 1000 mg/m²) orally twice a day D1-14, repeated at 3 weeks</p> <ul style="list-style-type: none"> · Drug: 5-fluorouracil/oxaliplatin <p>5-fluorouracil/oxaliplatin (FOLFOX): oxaliplatin 80 mg/m² (80% dose of 100 mg/m²) iv ov 120 min, leucovorin 80 mg/m² (80% dose of 100 mg/m²) iv ov 120min, 5-fluorouracil 1900 mg/m² (80% dose of 2400 mg/m²) iv ov 46h D1, repeated at 2 weeks</p> <p>Active Comparator: Fluoropyrimidine mono chemotherapy</p> <ul style="list-style-type: none"> · Drug: Capecitabine <p>Capecitabine : 1250 mg/m² orally twice a day D1-14 repeated at 3 weeks (if Ccr < 60 mL/min, 1000 mg/m² orally twice a day)</p> <ul style="list-style-type: none"> · Drug: S-1 <p>S-1 : 40 mg/m² orally twice a day D1-14 repeated at 3 weeks (if Ccr < 60 mL/min, 30 mg/m² orally twice a day)</p> <ul style="list-style-type: none"> · Drug: 5-fluorouracil <p>5-fluorouracil (FL) : leucovorin 100 mg/m² iv ov 2h, 5-fluorouracil 2400 mg/m² iv ov 46 h D1, repeated at 2 weeks</p> <p>Arm A (XP): cisplatin 50 mg/m² d1, capecitabine 1000 mg/m² orally twice a day d 1-14, repeated at day 21</p> <p>Arm B (SP): cisplatin 50mg/m² d1, S-1 30mg/m² orally twice a day d 1-14, repeated at d 21</p>

NCT02114359 (Continued)

Arm C (XELOX): oxaliplatin 100 mg/m²d1, capecitabine 800 mg/² orally twice a day d1 -14, repeated at day 21

Arm D (FOLFOX): oxaliplatin 80 mg/m², leucovorin 80 mg/m², 5-fluorouracil 1900 mg/m² ci 46 h d 1, repeated at day 14

Outcomes	Comparison of overall survival [Time frame: up to 3 years] Comparison of progression-free survival [Time frame: up to 2 years] Comparison of response rate [Time frame: up to 2 years] Comparison of adverse events [Time frame: up to 2 years] Comparison of quality of life [Time frame: up to 2 years]
Starting date	February 2014
Contact information	In Sil Choi, M.D., Ph.D.; 82-10-9137-3883; hmoischoi@hanmail.net
Notes	Sponsor and Collaborators: Seoul National University Hospital, Ministry of Health & Welfare, Korea, Korean Cancer Study Group

NCT02289378

Trial name or title	Dose-dense biweekly docetaxel, oxaliplatin and 5-fluorouracil as first-line treatment in advanced gastric cancer (DaeMon-Plus)
Methods	Phase 2 Intervention Model: Single-group assignment Masking: Open-label Primary Purpose: Treatment
Participants	N = 30 Age 18-75 years · Participants with histologically or cytologically confirmed unresectable gastric adenocarcinoma whose ECOG performance status are 0-2
Interventions	Experimental: docetaxel, oxaliplatin and 5-Fu Docetaxel 50 mg/m ² Oxaliplatin 85 mg/m ² 5-Fu 2800 mg/m ² Repeated every two weeks
Outcomes	Progression-free survival [Time frame: 2 years]
Starting date	November 2014
Contact information	Ping Lan, MD The Sixth Affiliated Hospital of Sun Yat-sen University
Notes	

NCT02289547

Trial name or title	Randomized phase 3 study of XELOX (capecitabine plus oxaliplatin) followed by maintenance capecitabine or observation in participants with advanced gastric adenocarcinoma
Methods	Multicentre RCT 2 arms Phase 3 Allocation: Randomised Intervention Model: Parallel assignment Masking: Open-label Primary Purpose: Treatment
Participants	N = 184 advanced or recurrent gastric cancer, age ≥18 years, HER-2 negative, with more than stable disease after 6 cycles 1st line of XELOX chemotherapy (objective response, non-complete response/non-progressive disease in cases of non-measurable disease before XELOX chemotherapy)
Interventions	Arm A : Capecitabine: capecitabine 1000 mg/m ² twice a day D1-14, repeated at 3 weeks Arm B : Observation
Outcomes	Progression-free survival, overall survival, quality of life (as measured by QLQ-c30 and STO-22), Toxicity profile
Starting date	January 2015
Contact information	Byoungyong Shim: shimby@catholic.ac.kr ; Ho Jung An:
Notes	Sponsor: The Catholic University of Korea

NCT02549911

Trial name or title	Hyperthermic intraperitoneal chemotherapy, intravenous chemotherapy combined with surgery for the treatment of advanced gastric cancer with peritoneal metastasis
Methods	Phase 2 Intervention Model: Single-group assignment Masking: Open-label Primary Purpose: Treatment
Participants	N = 40 18-75 years Gastric cancer confirmed by endoscopic biopsy , and enhanced CT suspected to have peritoneal metastasis, including ascites, ovarian metastasis, omentum or peritoneal metastasis.
Interventions	Experimental: HIPEC,Chemotherapy AND surgery 1. Surgical exploration,if PCI < 20,then we perform this study 2. HIPEC (RHL-2000B, Madain Medical Devices Co., Ltd., Jilin, China): Taxol (Paclitaxel Injection) 75 mg/m ² , twice, within 72 hours after surgical exploration ; oral chemotherapy:S-1(Tega-

NCT02549911 (Continued)

- fur,Gimeracil and Oteracil Potassium Capsules): 80 mg/m², twice daily (after the breakfast and supper) for two weeks, and then suspend for one week.
3. Chemotherapy (3 cycles) : Taxol 150 mg/m² ,d 1, S-1: 80 mg/m², twice daily (after breakfast and supper) for two weeks, and then suspend for one week.
 4. Surgery: Secondary surgical exploration:if PCI less than 20,then perform the cytoreductive surgery(resection of primary tumours and metastases)
 5. after the surgery,HIPEC for two cycles,and PS chemotherapy for 3 cycles
- Intervention: Other: HIPEC, chemotherapy AND surgery

Outcomes	R0 resection [Time frame: 3 months] Adverse events [Time frame: 6 months] Overall survival time [Time Frame: 3 years]
Starting date	September 2015
Contact information	Yian Du, MD; ypfzmu@163.com ; 86-571-88128031
Notes	

NCT02583659

Trial name or title	The first-line combined chemotherapy for advanced gastric cancer: A prospective observational clinical study
Methods	Observational Model: Cohort Time Perspective: Prospective
Participants	N = 250 Histopathology or cytopathology confirmed unresectable locally advanced, or recurrent, or metastatic chemotherapy-naive gastric cancer and gastroesophageal adenocarcinoma participants
Interventions	Observational Model: Cohort Time Perspective: Prospective
Outcomes	Overall survival (OS) [Time frame: From date of enrolment until the date of death from any cause, assessed up to 60 months] Progression-free survival (PFS) [Time frame: From date of enrolment until the date of first documented progression or date of death from any cause, whichever came first, assessed up to 60 months] Objective response rate (ORR) [Time frame: The sum of complete remission (CR) rate and partial remission (PR) rate. Response will be measured through first-line treatment completion, up to 1 year] Disease control rate (DCR) [Time frame: The sum of CR rate, PR rate and stable disease (SD) rate. Response will be measured through first-line treatment completion, up to 1 year] Number of participants with treatment-related adverse events as assessed by CTCAE v4.0 [Time frame: Through first-line treatment completion, up to 24 weeks.]
Starting date	January 2013

Chemotherapy for advanced gastric cancer (Review)

NCT02583659 (Continued)

Contact information Xianglin Yuan, MD,PHD
Tongji hospital of Tongji Medical College, Huazhong University of Science and Technology

Notes

NCT02855788

Trial name or title Phase II study of weekly metronomic chemotherapy using weekly aclitaxel, Oxaliplatin, Leucovorin and 5-FU (POLF) in participants with advanced gastric cancer

Methods Phase 2
Intervention Model: Single-group assignment
Masking: Open-label
Primary Purpose: Treatment

Participants N = 40
18-70 years
AJCC stage 3 or 4 gastric cancer

Interventions Experimental: POLF regimen
Paclitaxel 60 mg/m², oxaliplatin 50 mg/m², leucovorin 20 mg/m², and 5-FU 425 mg/m² IV weekly
Intervention: Drug: paclitaxel 60 mg/m², oxaliplatin 50 mg/m², leucovorin 20 mg/m², and 5-FU 425 mg/m² IV weekly

Outcomes Response rate [Time frame: 3 months]
-based on Recist 1.1
Adverse events [Time frame: 2 years]
-based on NCI-CTC v.2
Progression-free survival [Time frame: 2 years]
Overall survival [Time frame: 2 years]

Starting date May 2015

Contact information Contact: Nick N Chen, M.D., Ph.D. 206-588-1722 nicknchenmd@gmail.com
Contact: Jie Liu, M.D. 021-5288236 jieliu@fudan.edu.cn

Notes

NCT03006432

Trial name or title ASE III randomised trial to evaluate folfox with or without docetaxel (TFOX) as 1st line chemotherapy for locally advanced or metastatic oesophago-gastric carcinoma (GASTFOX)

NCT03006432 (Continued)

Methods	Phase 3 Allocation: Randomised Intervention Model: Parallel assignment Masking: No masking Primary Purpose: Treatment
Participants	N = 506 Gastric or gastro-oesophageal junction adenocarcinoma (all Siewert), histologically proven (on primary tumour or metastatic lesion), HER2 negative (positive HER2 status is defined by a positive IHC test of 3+ or IHC of 2+ with positive FISH) Metastatic or non-resectable (locally advanced) disease
Interventions	Active Comparator: FOLFOX Cycles every 15 days until progression disease Experimental: TFOX Cycles every 15 days until progression disease Interventions: Drug: oxaliplatin 1. Drug: 5Fluorouracil bolus 2. Drug: 5Fluorouracil continuous 3. Drug: docetaxel 4. Drug: folinic Acid
Outcomes	Progression-free survival [Time frame: 12 months after last randomisation] Overall survival toxicity events (adverse events) according to NCI-CTC v4.0 [Time frame: 12 months after last randomisation] Objective response rate [Time frame: 12 months after last randomisation] Toxicity events according to NCI-CTC v4.0 [Time frame: 12 months after last randomisation]
Starting date	December 2016
Contact information	Contact: Marie MOREAU +33 (0)380393404 marie.moreau@u-bourgogne.fr
Notes	PRODIGE 51

Tsuburaya 2012

Trial name or title	A randomized phase II trial to elucidate the efficacy of capecitabine plus cisplatin (XP) and S-1 plus cisplatin (SP) as a first-line treatment for advanced gastric cancer: XP ascertainment vs. SP randomized PII trial (XParTS II)
Methods	Multicentre RCT 2 arms

Tsuburaya 2012 (Continued)

Participants	N = 100 (planned)
Interventions	Arm A: S-1 (40 mg/m ² twice a day d1-21) + cisplatin (60 mg/m ² d 8) repeated at d 35 versus Arm B: capecitabine (1000 mg/m ² d1-14) + cisplatin (80 mg/m ² d 1) repeated at d 21
Outcomes	Progression-free survival, overall survival, time to treatment failure, tumour response, safety
Starting date	August 2011
Contact information	Akira Tsuburaya, Department of Gastrointestinal Surgery, Kanagawa Cancer Center, 1-1-2, Nakao, 241-0815, Yokohama, Asahi-ku, Japan, tuburayaa@kcch.jp
Notes	The study is registered (NCT01406249), the study protocol is published. Estimated study completion date is June 2015. The study is not yet published.

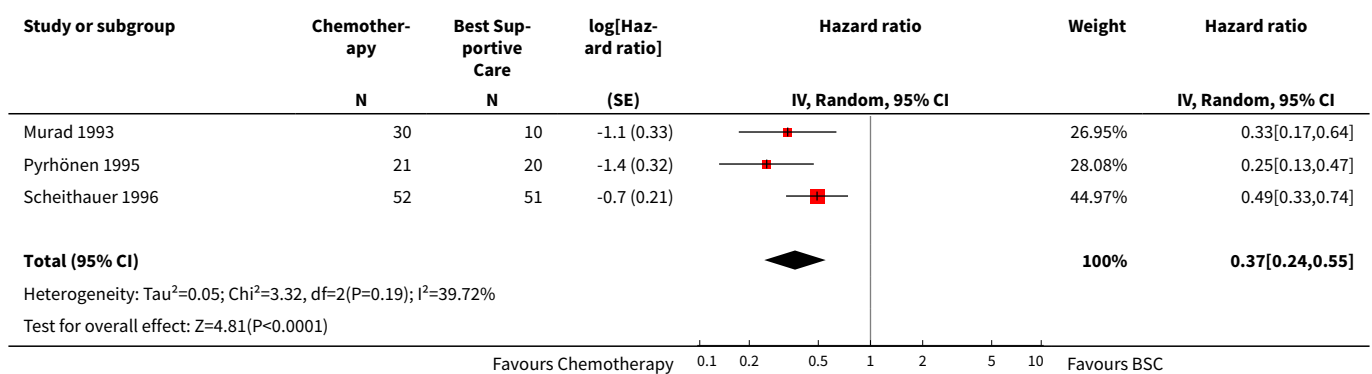
5-FU: 5-fluorouracil
ci: continuous infusion
d: day
E: epirubicin
G-CSF: granulocyte colony stimulating factor
PFS: progression-free survival
RCT: randomised controlled trial

DATA AND ANALYSES

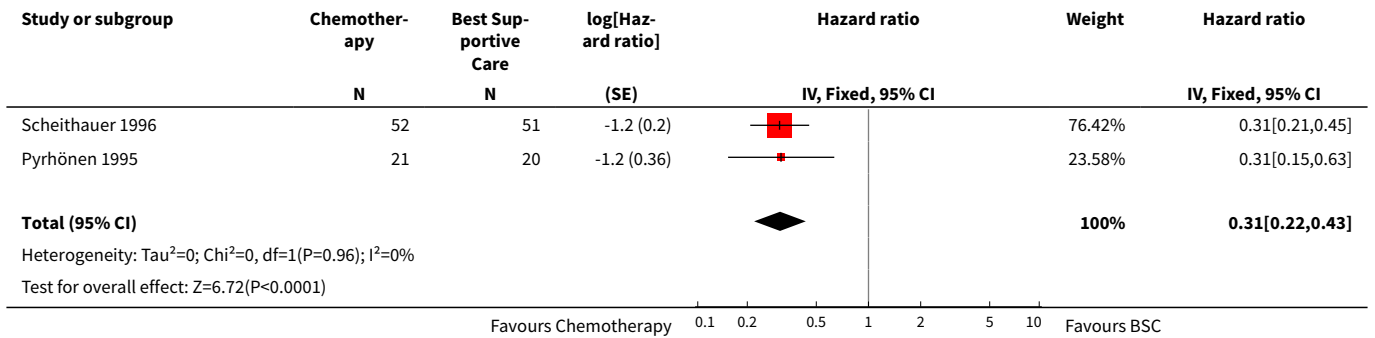
Comparison 1. Chemotherapy versus best supportive care

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Overall survival	3	184	Hazard ratio (Random, 95% CI)	0.37 [0.24, 0.55]
2 Time to progression	2	144	Hazard ratio (Fixed, 95% CI)	0.31 [0.22, 0.43]

Analysis 1.1. Comparison 1 Chemotherapy versus best supportive care, Outcome 1 Overall survival.



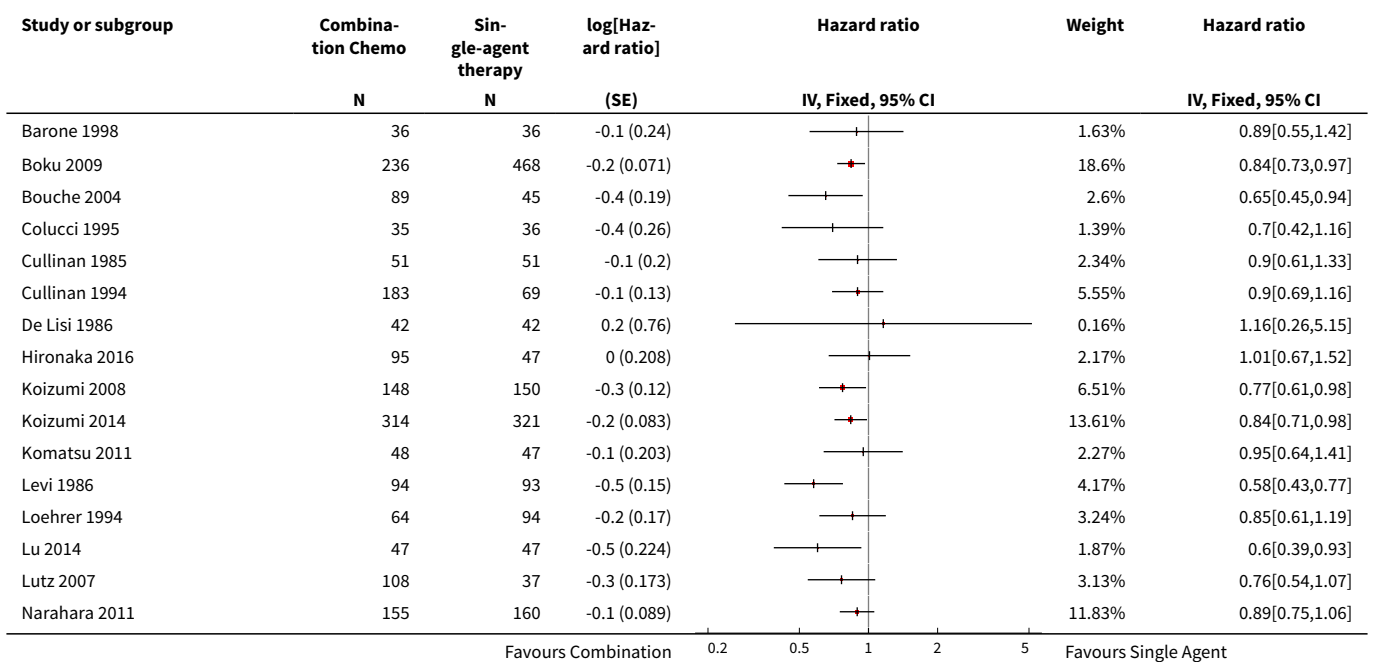
Analysis 1.2. Comparison 1 Chemotherapy versus best supportive care, Outcome 2 Time to progression.

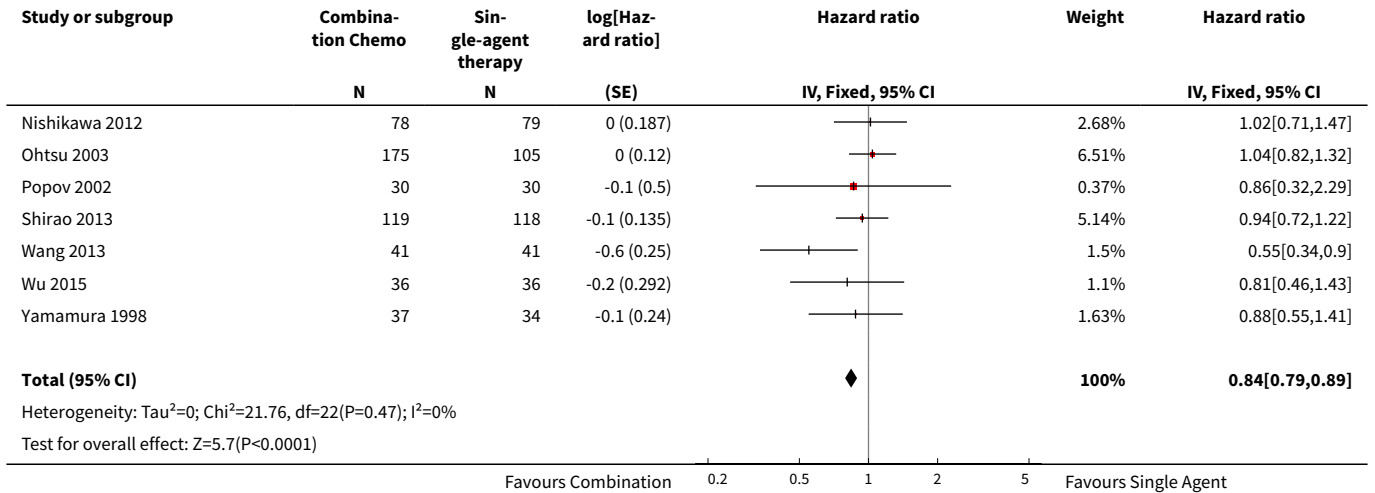


Comparison 2. Combination versus single-agent chemotherapy

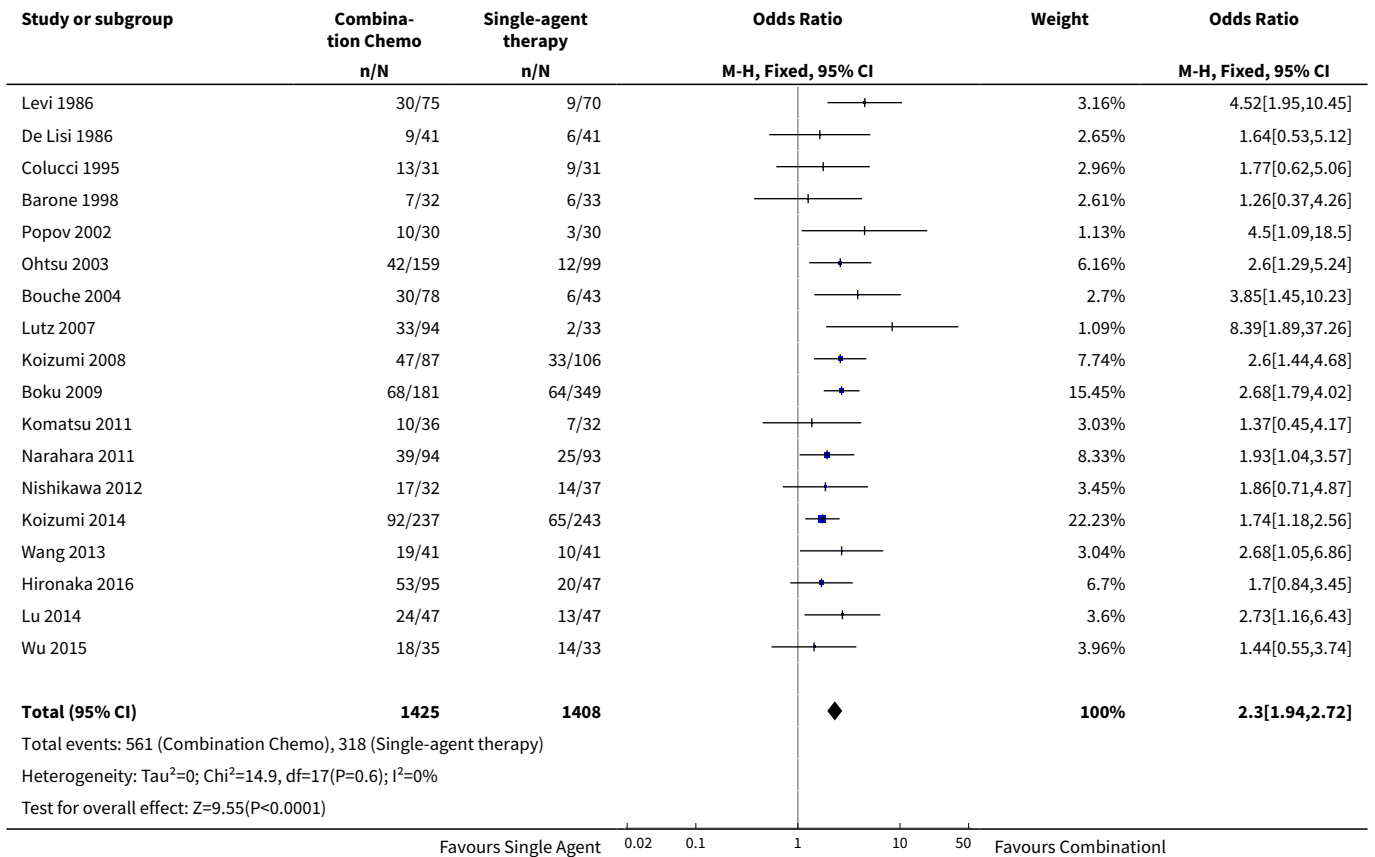
Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Overall survival	23	4447	Hazard ratio (Fixed, 95% CI)	0.84 [0.79, 0.89]
2 Tumour response	18	2833	Odds Ratio (M-H, Fixed, 95% CI)	2.30 [1.94, 2.72]
3 Time to progression	4	720	Hazard ratio (Random, 95% CI)	0.69 [0.55, 0.87]
4 Treatment-related death	18	3876	Odds Ratio (M-H, Fixed, 95% CI)	1.64 [0.83, 3.24]

Analysis 2.1. Comparison 2 Combination versus single-agent chemotherapy, Outcome 1 Overall survival.

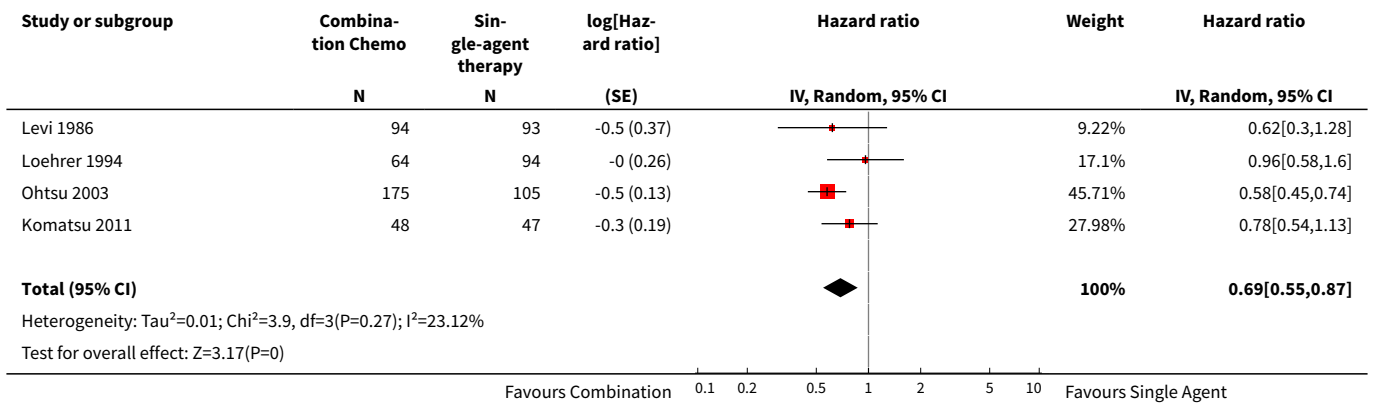




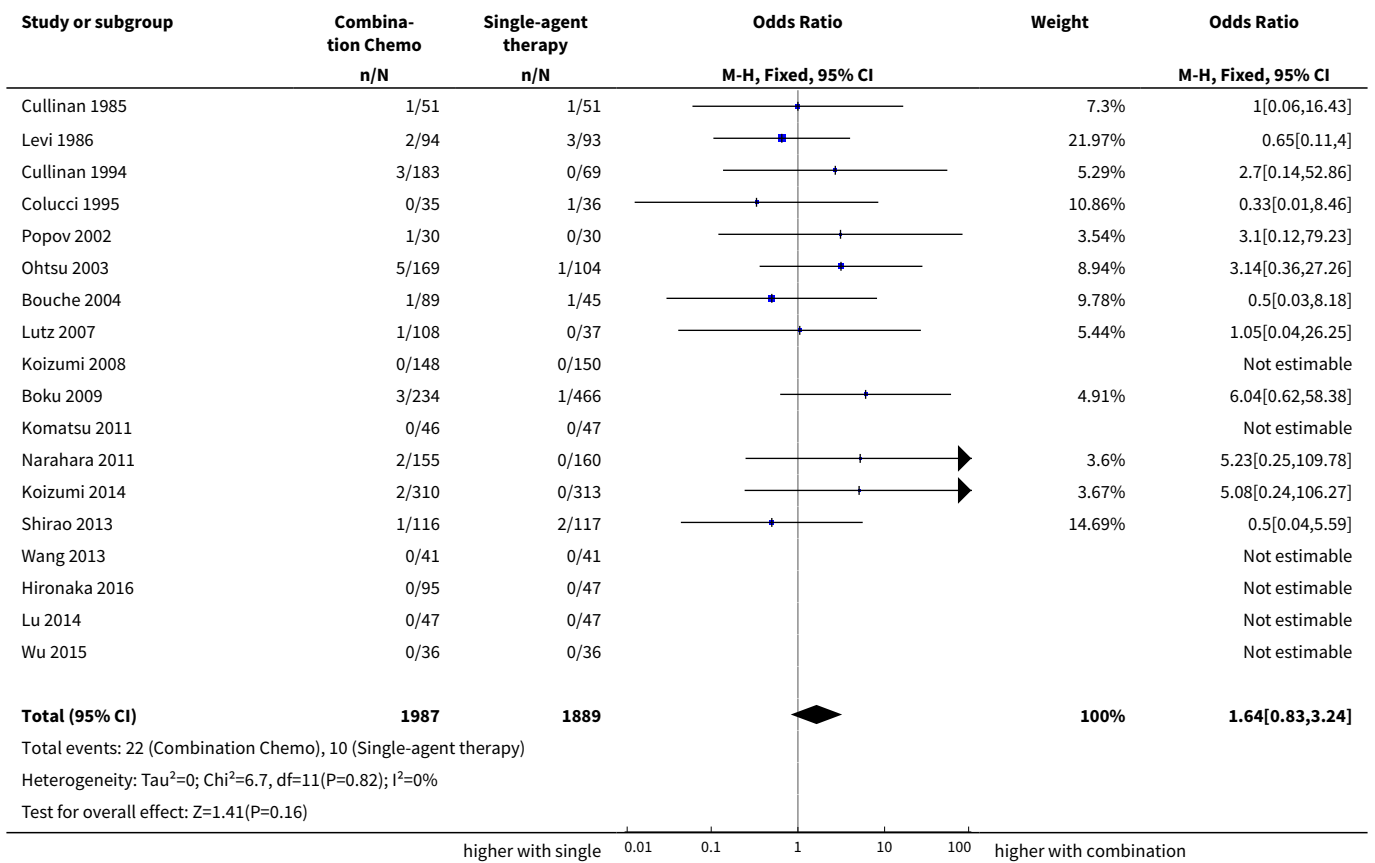
Analysis 2.2. Comparison 2 Combination versus single-agent chemotherapy, Outcome 2 Tumour response.



Analysis 2.3. Comparison 2 Combination versus single-agent chemotherapy, Outcome 3 Time to progression.



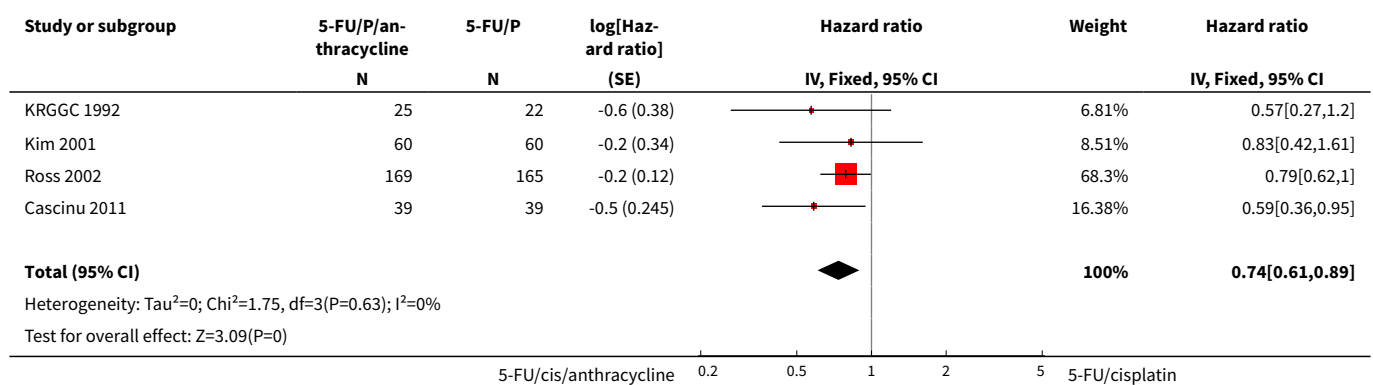
Analysis 2.4. Comparison 2 Combination versus single-agent chemotherapy, Outcome 4 Treatment-related death.



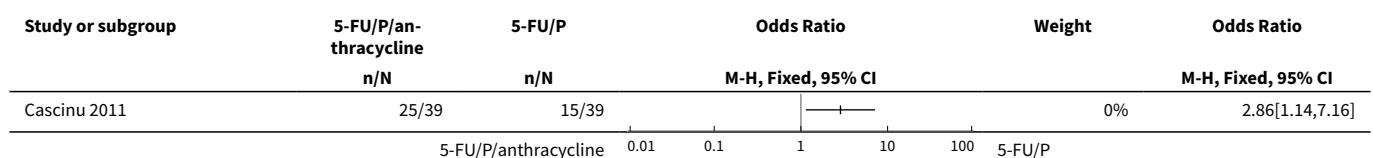
Comparison 3. 5-FU/cisplatin/anthracycline combinations versus 5-FU/cisplatin combinations (without anthracyclines)

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Overall survival	4	579	Hazard ratio (Fixed, 95% CI)	0.74 [0.61, 0.89]
2 Tumour response	1		Odds Ratio (M-H, Fixed, 95% CI)	Subtotals only
3 Time to progression	1		Hazard Ratio (Fixed, 95% CI)	Subtotals only

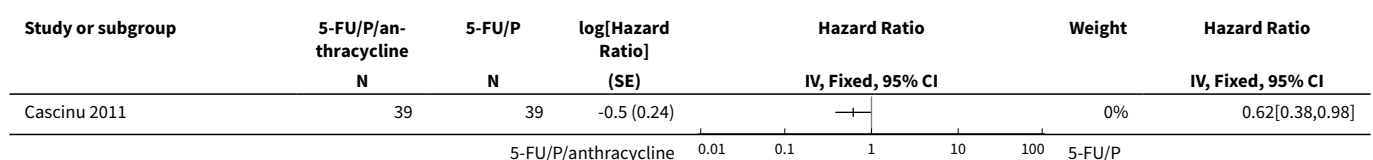
Analysis 3.1. Comparison 3 5-FU/cisplatin/anthracycline combinations versus 5-FU/cisplatin combinations (without anthracyclines), Outcome 1 Overall survival.



Analysis 3.2. Comparison 3 5-FU/cisplatin/anthracycline combinations versus 5-FU/cisplatin combinations (without anthracyclines), Outcome 2 Tumour response.



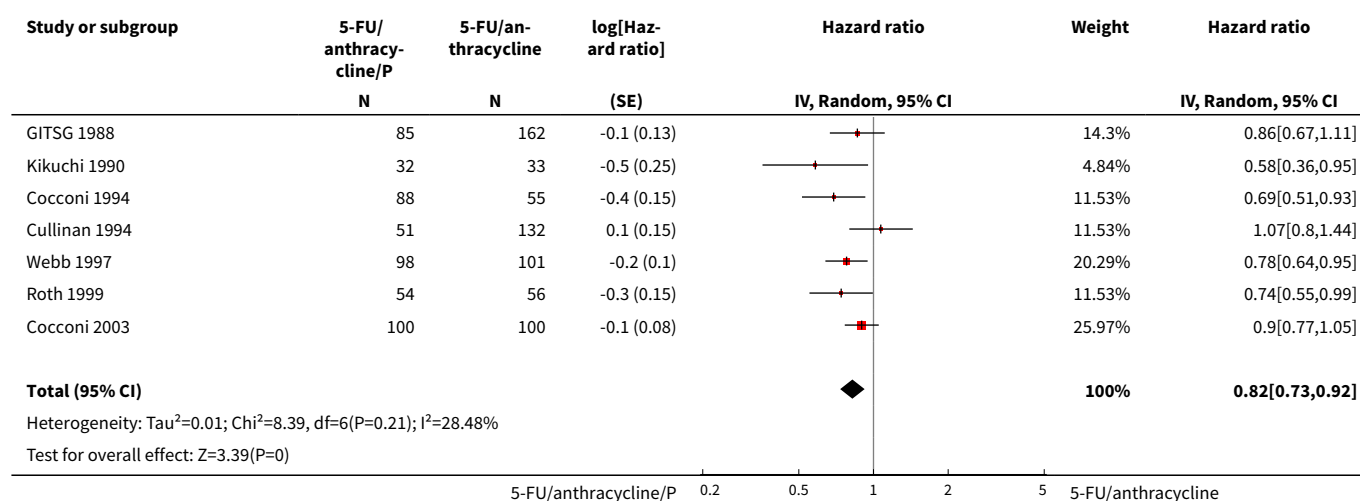
Analysis 3.3. Comparison 3 5-FU/cisplatin/anthracycline combinations versus 5-FU/cisplatin combinations (without anthracyclines), Outcome 3 Time to progression.



Comparison 4. 5-FU/cisplatin/anthracycline combinations versus 5-FU/anthracycline combinations (without cisplatin)

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Overall survival	7	1147	Hazard ratio (Random, 95% CI)	0.82 [0.73, 0.92]

Analysis 4.1. Comparison 4 5-FU/cisplatin/anthracycline combinations versus 5-FU/anthracycline combinations (without cisplatin), Outcome 1 Overall survival.

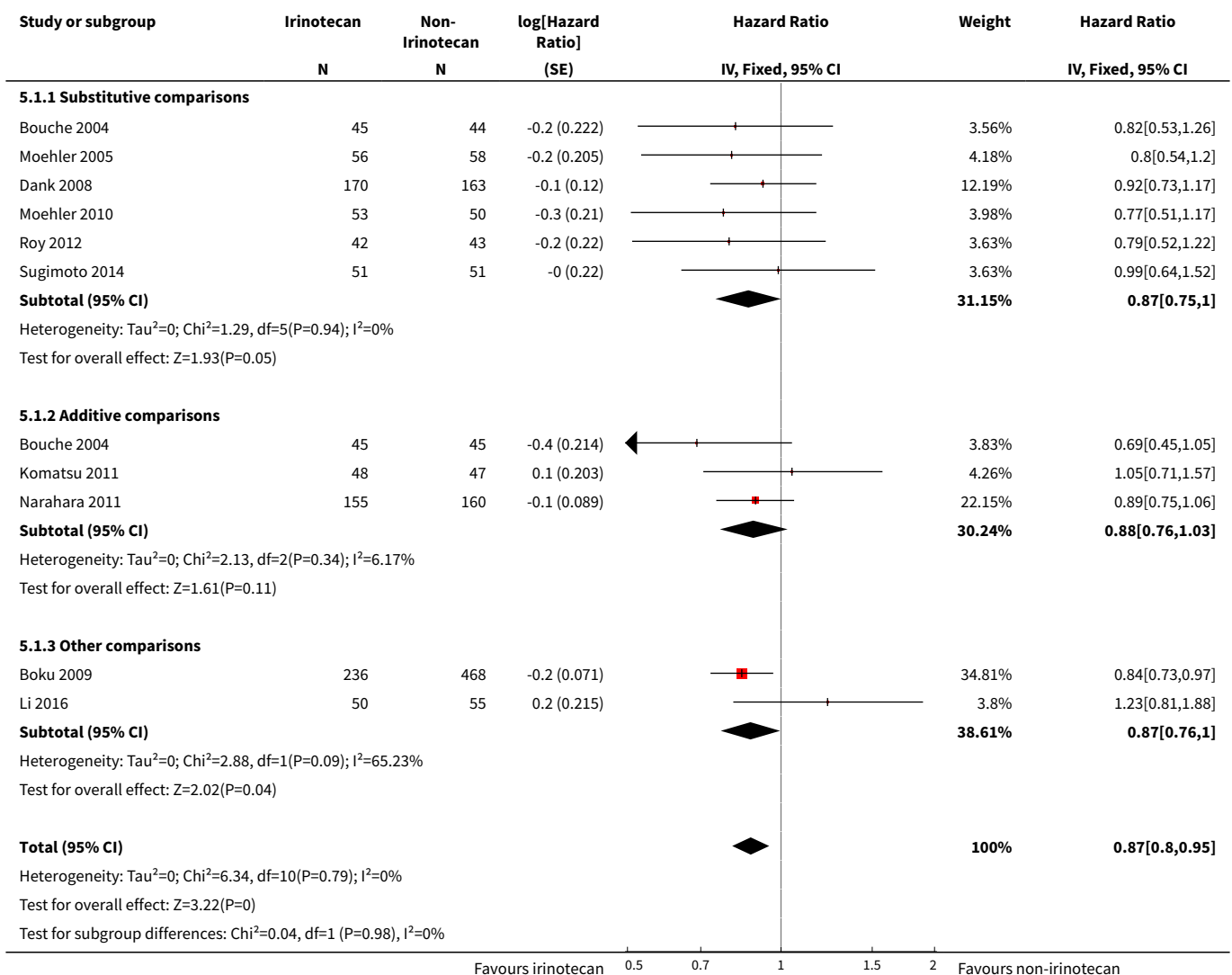


Comparison 5. Chemotherapy with irinotecan versus non-irinotecan-containing regimes

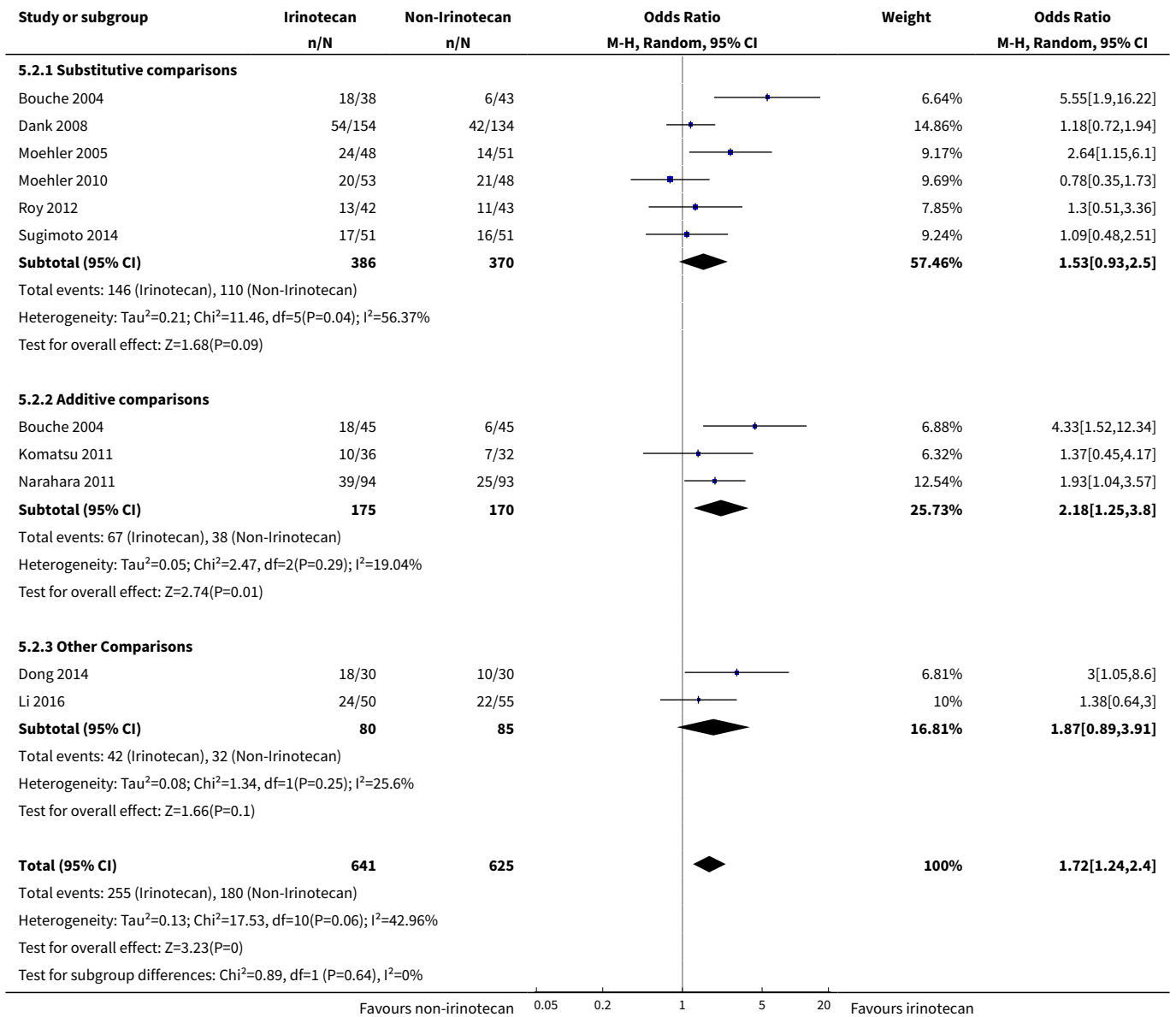
Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Overall survival	10	2135	Hazard Ratio (Fixed, 95% CI)	0.87 [0.80, 0.95]
1.1 Substitutive comparisons	6	826	Hazard Ratio (Fixed, 95% CI)	0.87 [0.75, 1.00]
1.2 Additive comparisons	3	500	Hazard Ratio (Fixed, 95% CI)	0.88 [0.76, 1.03]
1.3 Other comparisons	2	809	Hazard Ratio (Fixed, 95% CI)	0.87 [0.76, 1.00]
2 Tumour response	10	1266	Odds Ratio (M-H, Random, 95% CI)	1.72 [1.24, 2.40]
2.1 Substitutive comparisons	6	756	Odds Ratio (M-H, Random, 95% CI)	1.53 [0.93, 2.50]
2.2 Additive comparisons	3	345	Odds Ratio (M-H, Random, 95% CI)	2.18 [1.25, 3.80]
2.3 Other Comparisons	2	165	Odds Ratio (M-H, Random, 95% CI)	1.87 [0.89, 3.91]
3 Progression-free survival	7	1640	Hazard Ratio (Fixed, 95% CI)	0.76 [0.69, 0.84]
3.1 Substitutive comparison	5	741	Hazard Ratio (Fixed, 95% CI)	0.85 [0.72, 1.00]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
3.2 Additive comparisons	1	90	Hazard Ratio (Fixed, 95% CI)	0.51 [0.33, 0.77]
3.3 Other comparisons	2	809	Hazard Ratio (Fixed, 95% CI)	0.74 [0.66, 0.84]
4 Treatment-related death	9	1979	Odds Ratio (M-H, Random, 95% CI)	0.88 [0.23, 3.32]
5 Treatment discontinuation due to toxicity	9	1979	Odds Ratio (M-H, Random, 95% CI)	1.00 [0.46, 2.20]

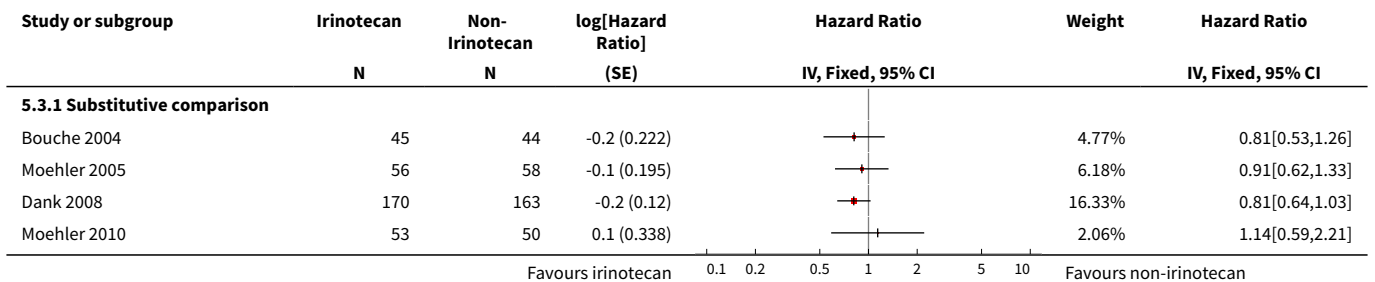
Analysis 5.1. Comparison 5 Chemotherapy with irinotecan versus non-irinotecan-containing regimes, Outcome 1 Overall survival.

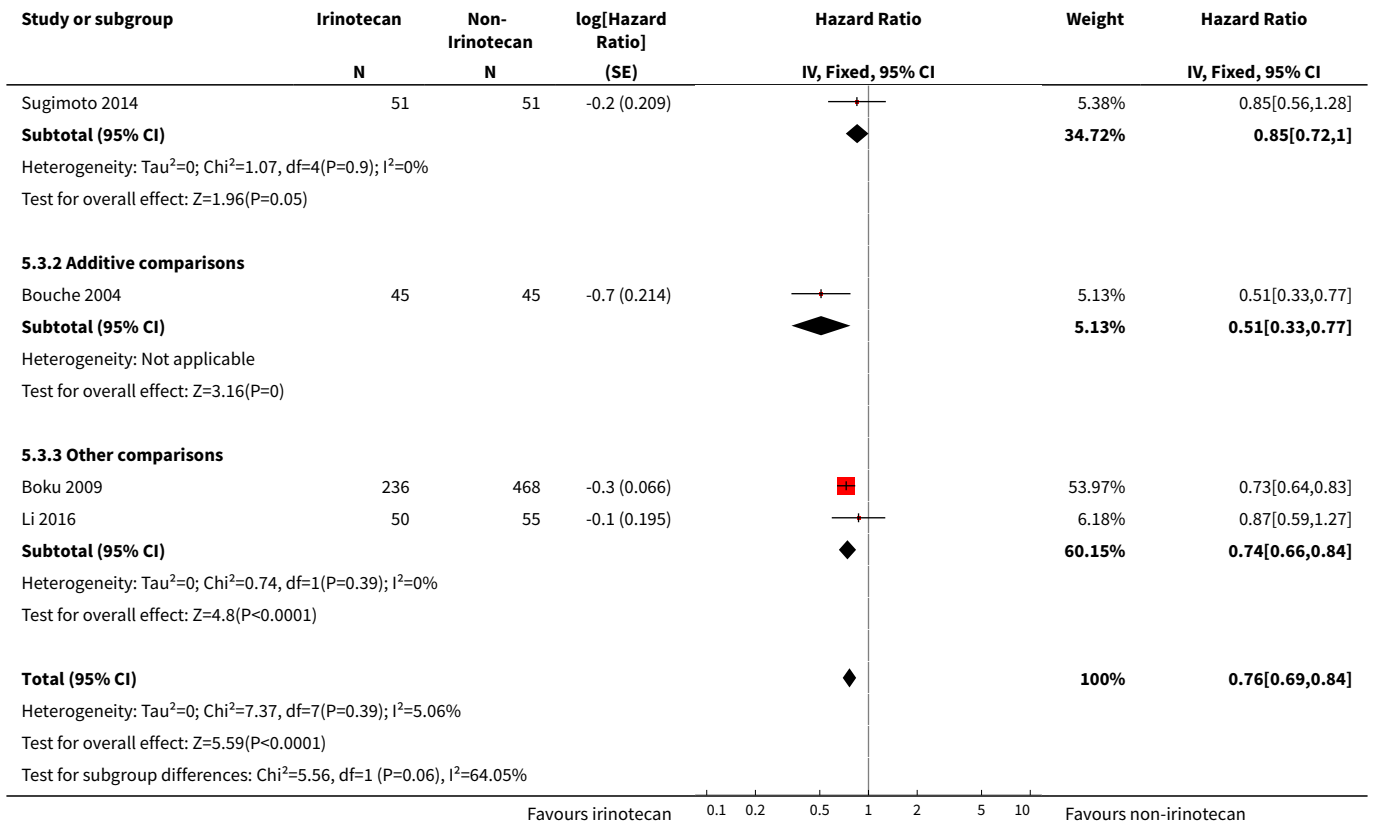


Analysis 5.2. Comparison 5 Chemotherapy with irinotecan versus non-irinotecan-containing regimes, Outcome 2 Tumour response.

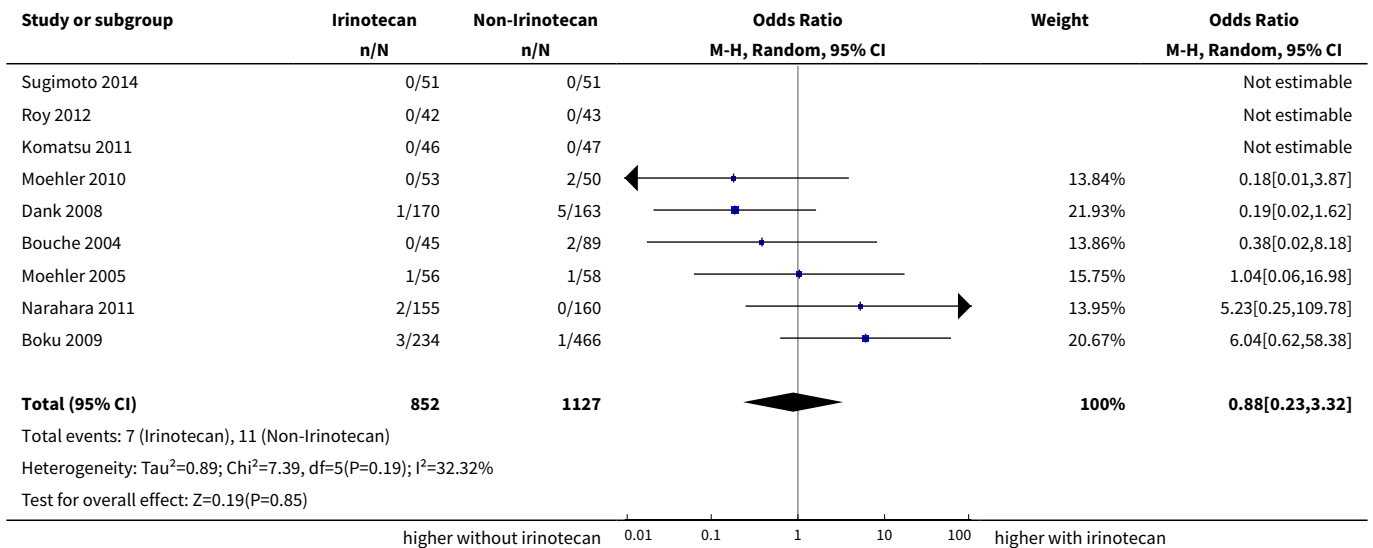


Analysis 5.3. Comparison 5 Chemotherapy with irinotecan versus non-irinotecan-containing regimes, Outcome 3 Progression-free survival.

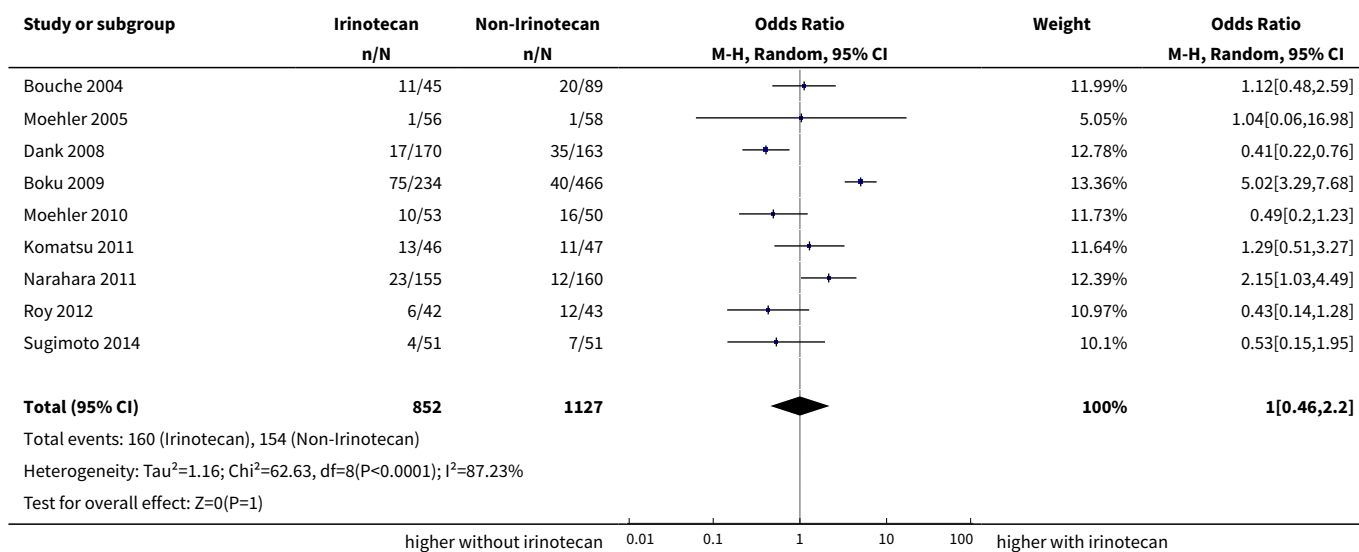




Analysis 5.4. Comparison 5 Chemotherapy with irinotecan versus non-irinotecan-containing regimes, Outcome 4 Treatment-related death.



Analysis 5.5. Comparison 5 Chemotherapy with irinotecan versus non-irinotecan-containing regimes, Outcome 5 Treatment discontinuation due to toxicity.

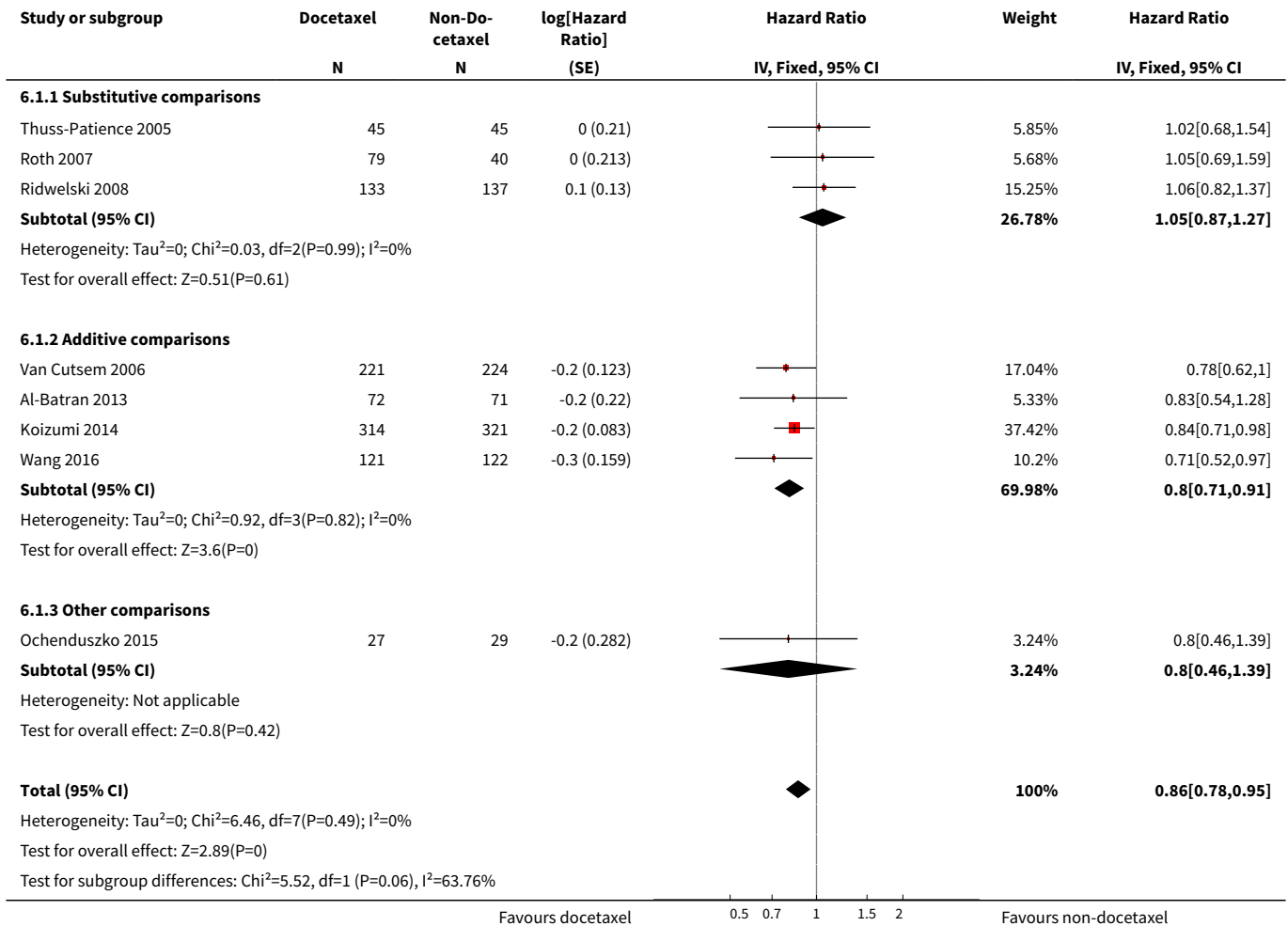


Comparison 6. Chemotherapy with docetaxel versus non-docetaxel-containing regimes

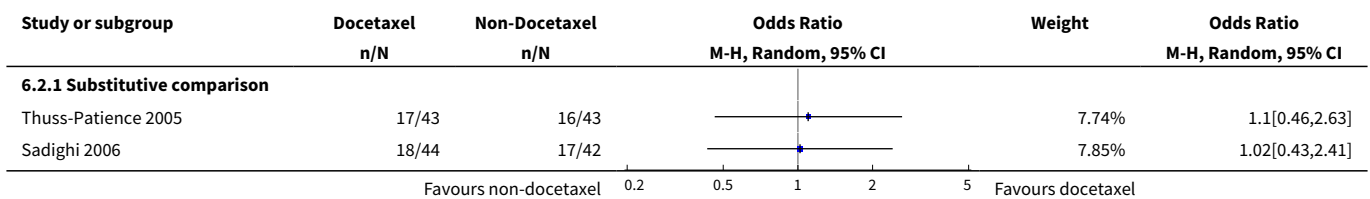
Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Overall survival	8	2001	Hazard Ratio (Fixed, 95% CI)	0.86 [0.78, 0.95]
1.1 Substitutive comparisons	3	479	Hazard Ratio (Fixed, 95% CI)	1.05 [0.87, 1.27]
1.2 Additive comparisons	4	1466	Hazard Ratio (Fixed, 95% CI)	0.80 [0.71, 0.91]
1.3 Other comparisons	1	56	Hazard Ratio (Fixed, 95% CI)	0.80 [0.46, 1.39]
2 Tumour response	9	1820	Odds Ratio (M-H, Random, 95% CI)	1.37 [1.03, 1.83]
2.1 Substitutive comparison	4	525	Odds Ratio (M-H, Random, 95% CI)	1.03 [0.71, 1.50]
2.2 Additive comparison	4	1235	Odds Ratio (M-H, Random, 95% CI)	1.83 [1.45, 2.32]
2.3 Other comparisons	1	60	Odds Ratio (M-H, Random, 95% CI)	0.33 [0.12, 0.96]
3 Time to progression	2	360	Hazard Ratio (Random, 95% CI)	1.06 [0.85, 1.32]
4 Progression-free survival	5	1498	Hazard Ratio (Random, 95% CI)	0.76 [0.63, 0.91]
4.1 Substitutive comparisons	1	119	Hazard Ratio (Random, 95% CI)	1.15 [0.77, 1.72]
4.2 Additive comparison (PFS)	3	1323	Hazard Ratio (Random, 95% CI)	0.70 [0.61, 0.81]
4.3 Other comparisons	1	56	Hazard Ratio (Random, 95% CI)	0.94 [0.55, 1.60]
5 Treatment-related death	7	2113	Odds Ratio (M-H, Fixed, 95% CI)	1.10 [0.55, 2.20]

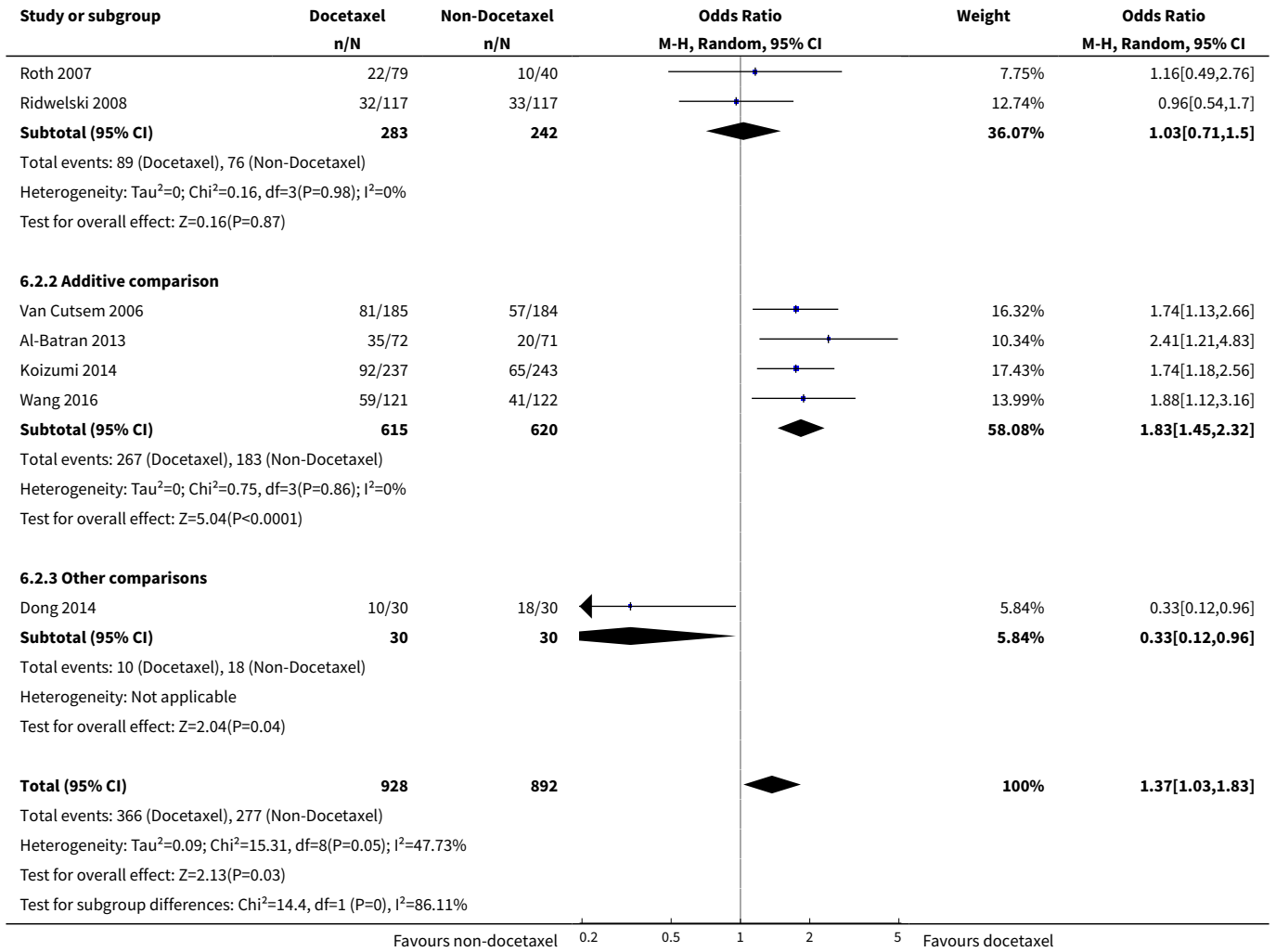
Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
6 Treatment discontinuation due to toxicity	5	1066	Odds Ratio (M-H, Random, 95% CI)	0.81 [0.53, 1.25]

Analysis 6.1. Comparison 6 Chemotherapy with docetaxel versus non-docetaxel-containing regimes, Outcome 1 Overall survival.

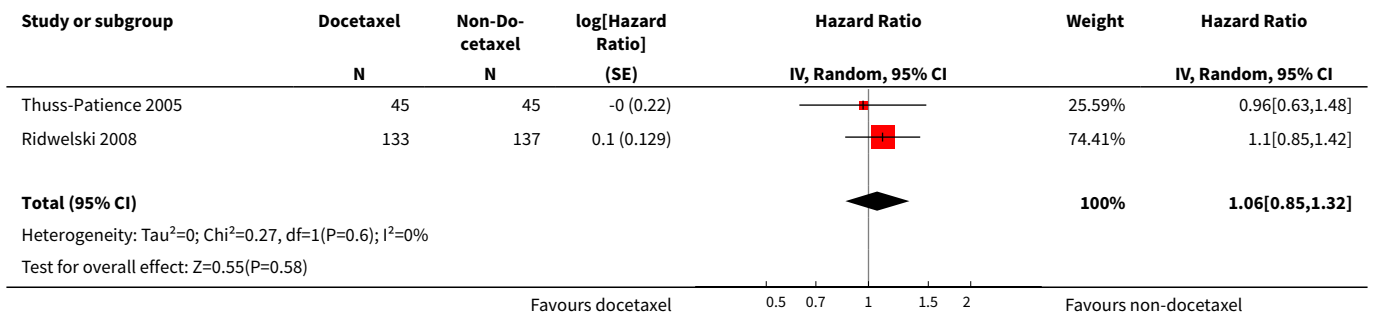


Analysis 6.2. Comparison 6 Chemotherapy with docetaxel versus non-docetaxel-containing regimes, Outcome 2 Tumour response.

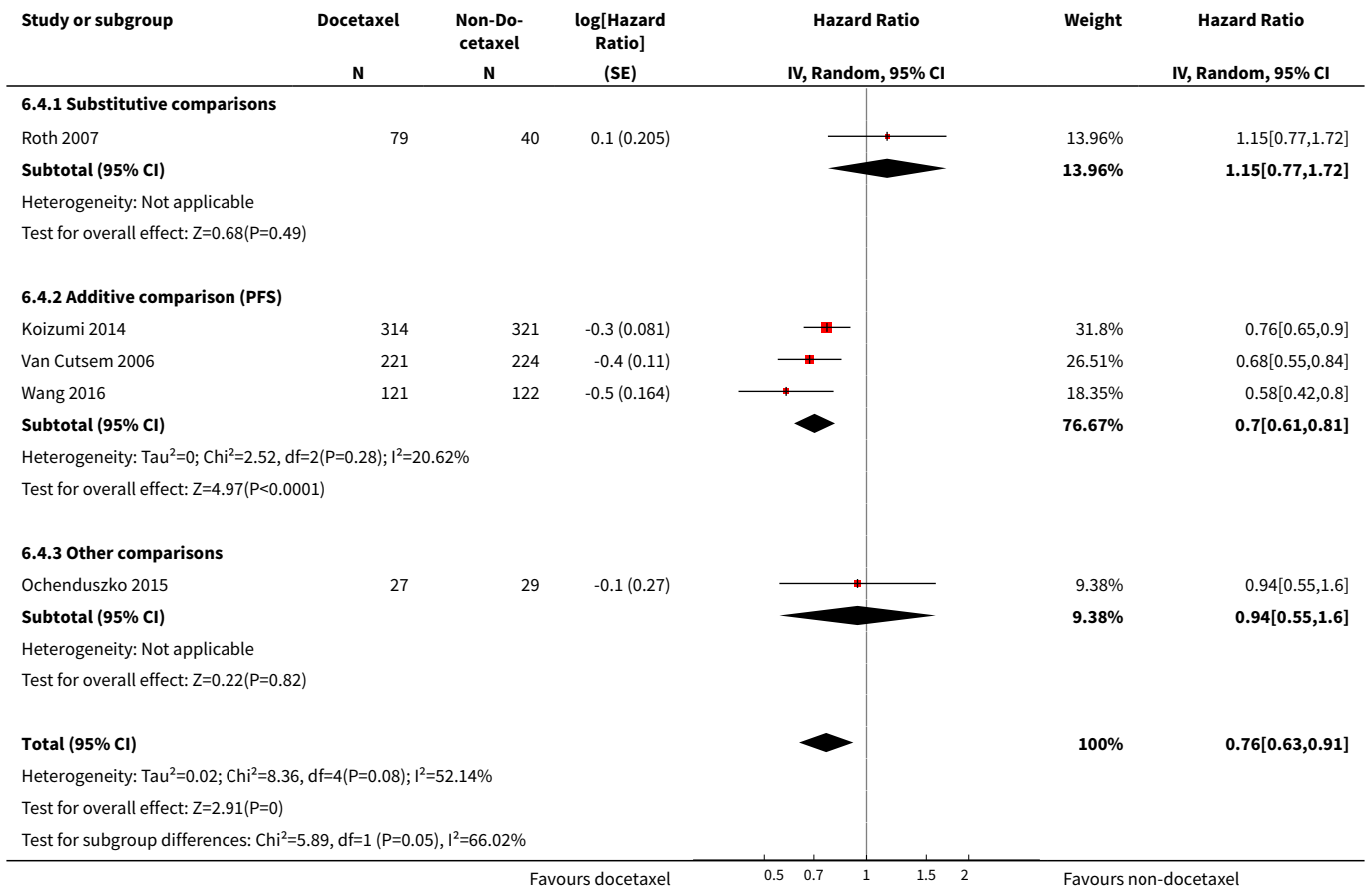




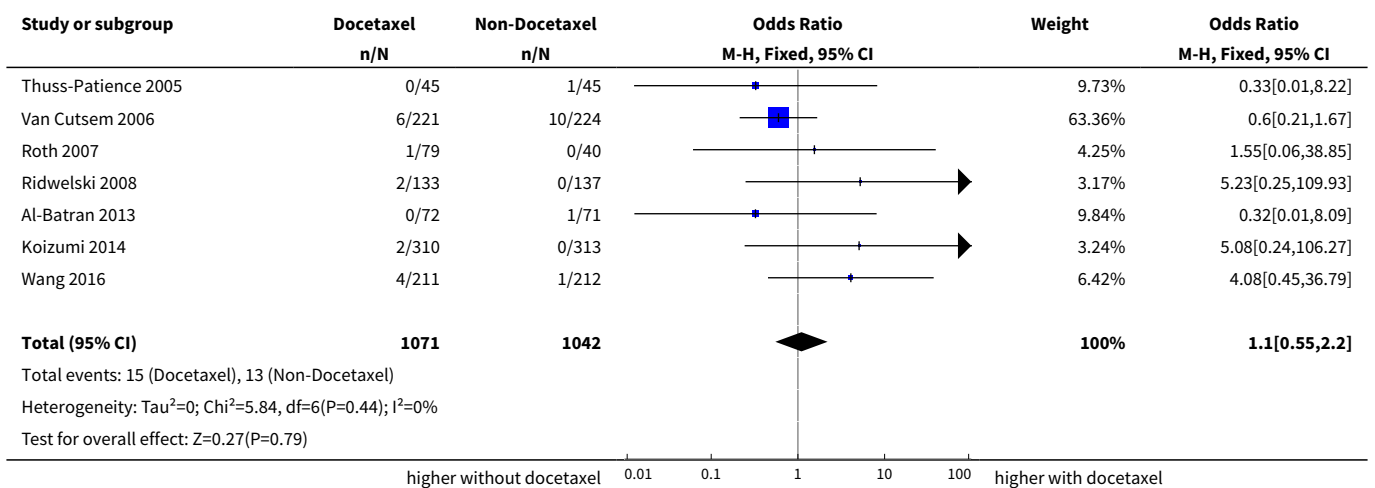
Analysis 6.3. Comparison 6 Chemotherapy with docetaxel versus non-docetaxel-containing regimes, Outcome 3 Time to progression.



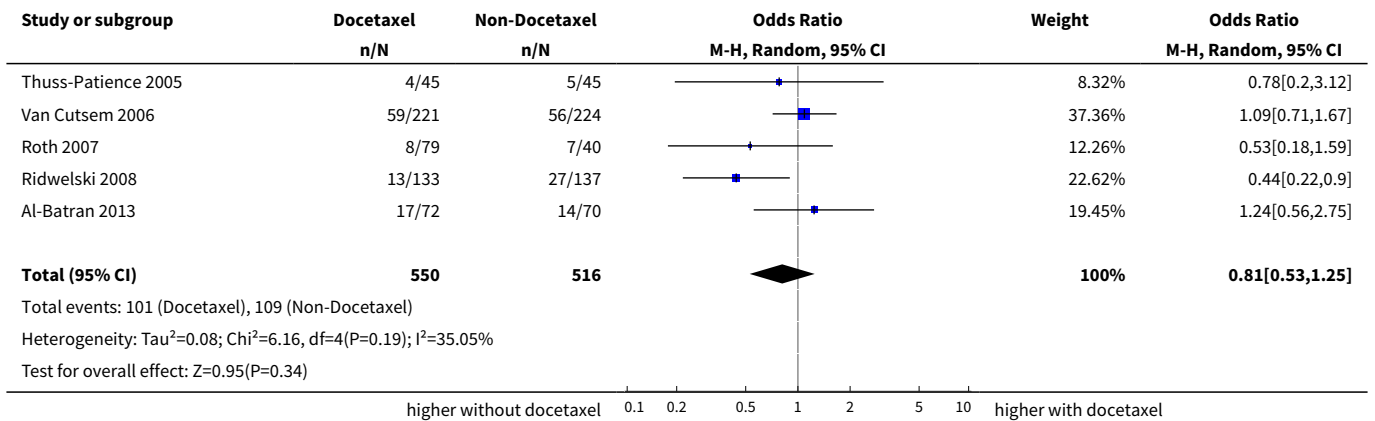
Analysis 6.4. Comparison 6 Chemotherapy with docetaxel versus non-docetaxel-containing regimes, Outcome 4 Progression-free survival.



Analysis 6.5. Comparison 6 Chemotherapy with docetaxel versus non-docetaxel-containing regimes, Outcome 5 Treatment-related death.



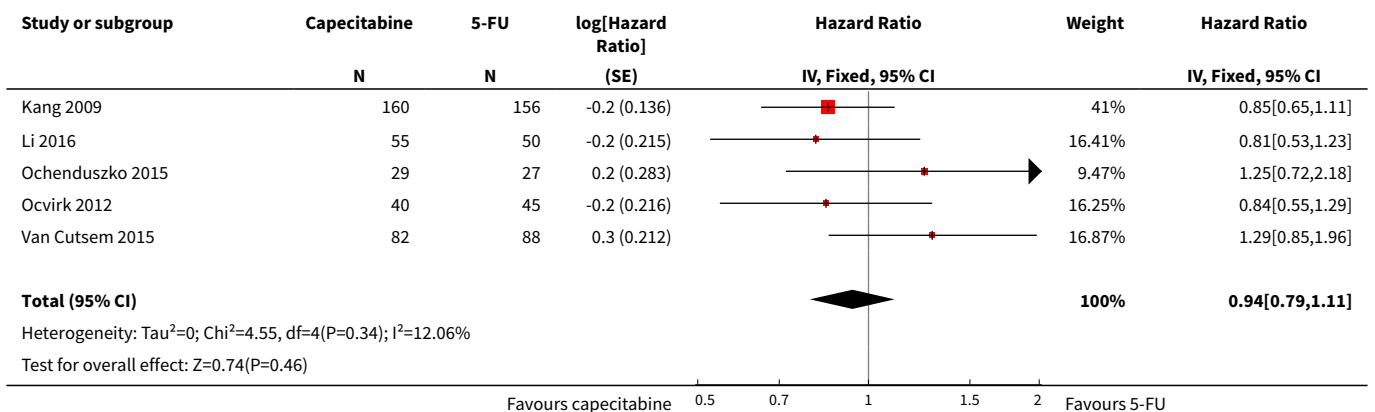
Analysis 6.6. Comparison 6 Chemotherapy with docetaxel versus non-docetaxel-containing regimes, Outcome 6 Treatment discontinuation due to toxicity.



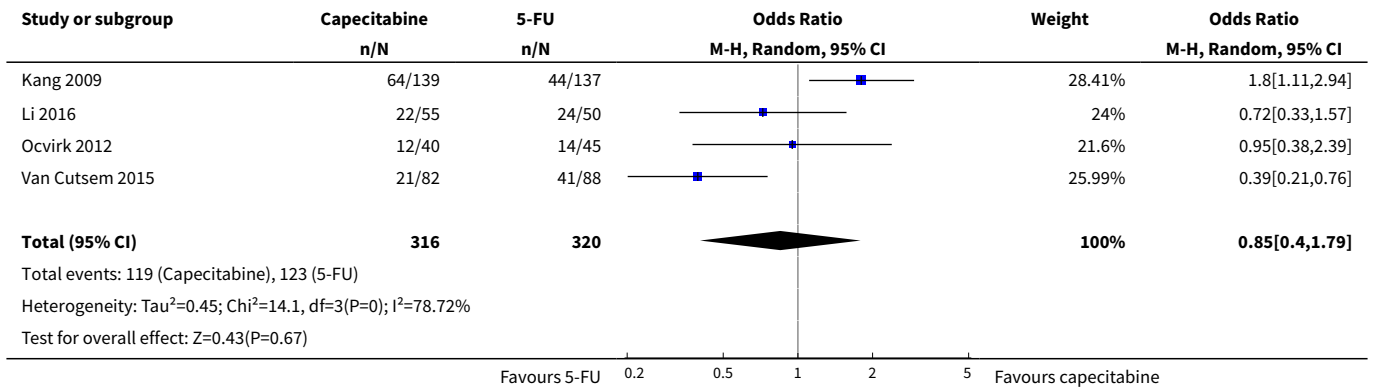
Comparison 7. Chemotherapy with capecitabine versus 5-FU-containing regimes

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Overall Survival	5	732	Hazard Ratio (Fixed, 95% CI)	0.94 [0.79, 1.11]
2 Tumour response	4	636	Odds Ratio (M-H, Random, 95% CI)	0.85 [0.40, 1.79]
3 Time to progression	1		Hazard Ratio (Fixed, 95% CI)	Subtotals only
4 Progression-free survival	4	647	Hazard Ratio (Random, 95% CI)	0.98 [0.77, 1.23]
5 Treatment-related death	2	481	Odds Ratio (M-H, Random, 95% CI)	1.88 [0.23, 15.15]
6 Treatment discontinuation due to toxicity	1		Odds Ratio (M-H, Fixed, 95% CI)	Subtotals only

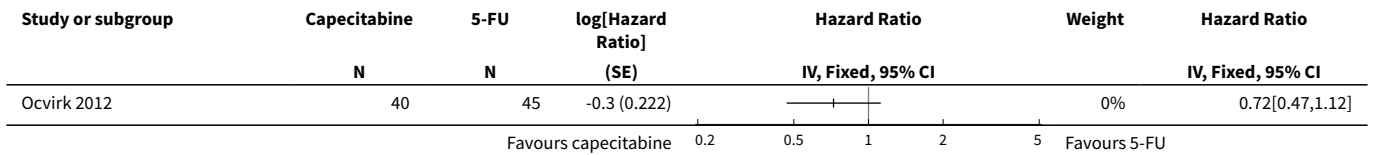
Analysis 7.1. Comparison 7 Chemotherapy with capecitabine versus 5-FU-containing regimes, Outcome 1 Overall Survival.



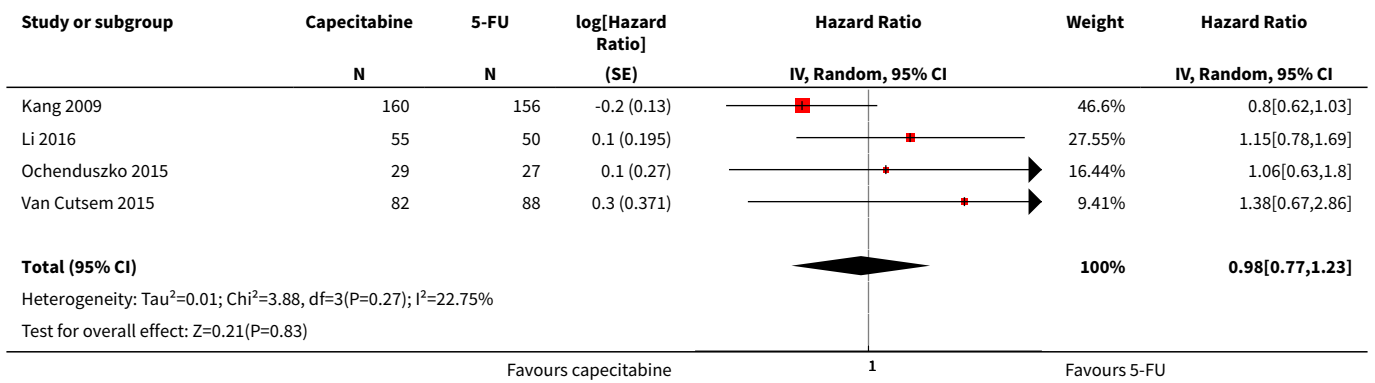
Analysis 7.2. Comparison 7 Chemotherapy with capecitabine versus 5-FU-containing regimes, Outcome 2 Tumour response.



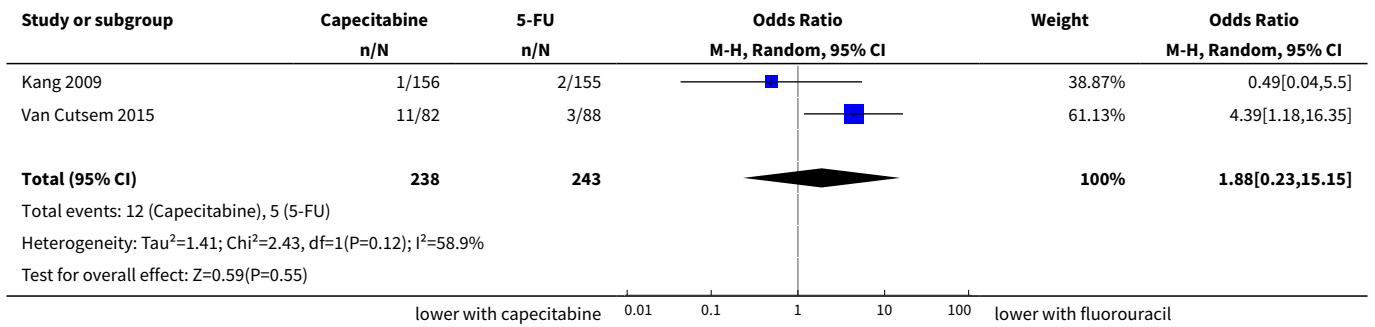
Analysis 7.3. Comparison 7 Chemotherapy with capecitabine versus 5-FU-containing regimes, Outcome 3 Time to progression.



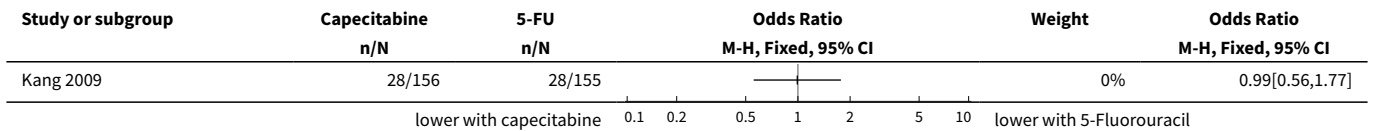
Analysis 7.4. Comparison 7 Chemotherapy with capecitabine versus 5-FU-containing regimes, Outcome 4 Progression-free survival.



Analysis 7.5. Comparison 7 Chemotherapy with capecitabine versus 5-FU-containing regimens, Outcome 5 Treatment-related death.



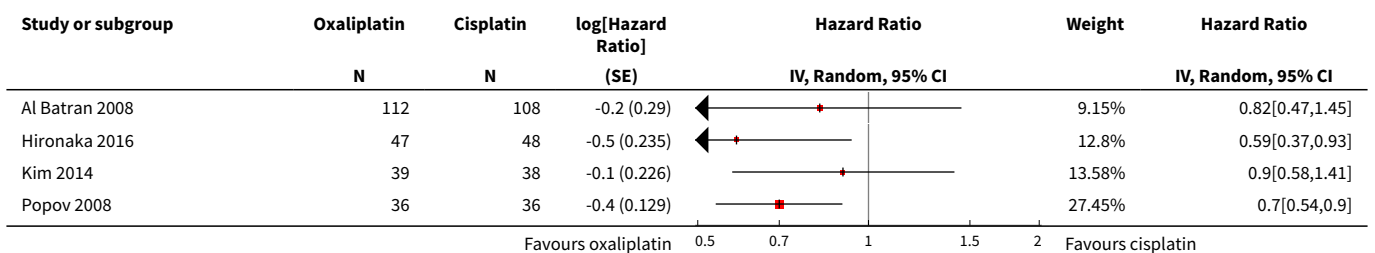
Analysis 7.6. Comparison 7 Chemotherapy with capecitabine versus 5-FU-containing regimens, Outcome 6 Treatment discontinuation due to toxicity.

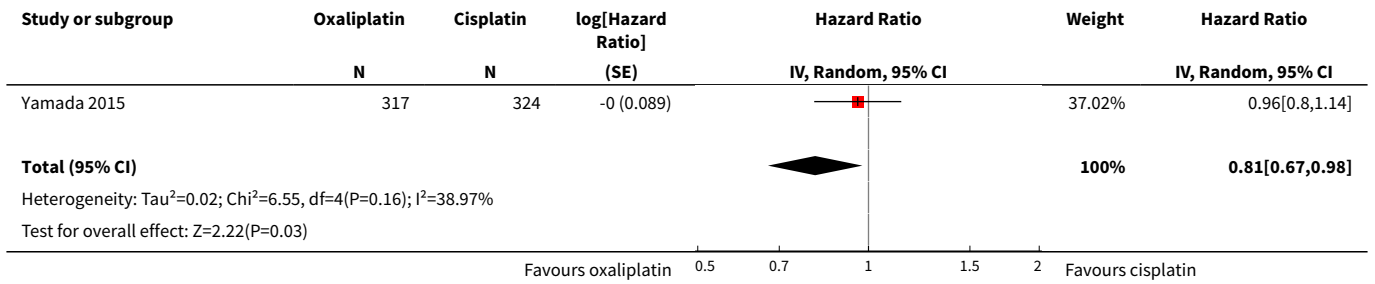


Comparison 8. Chemotherapy with oxaliplatin versus the same regime including cisplatin

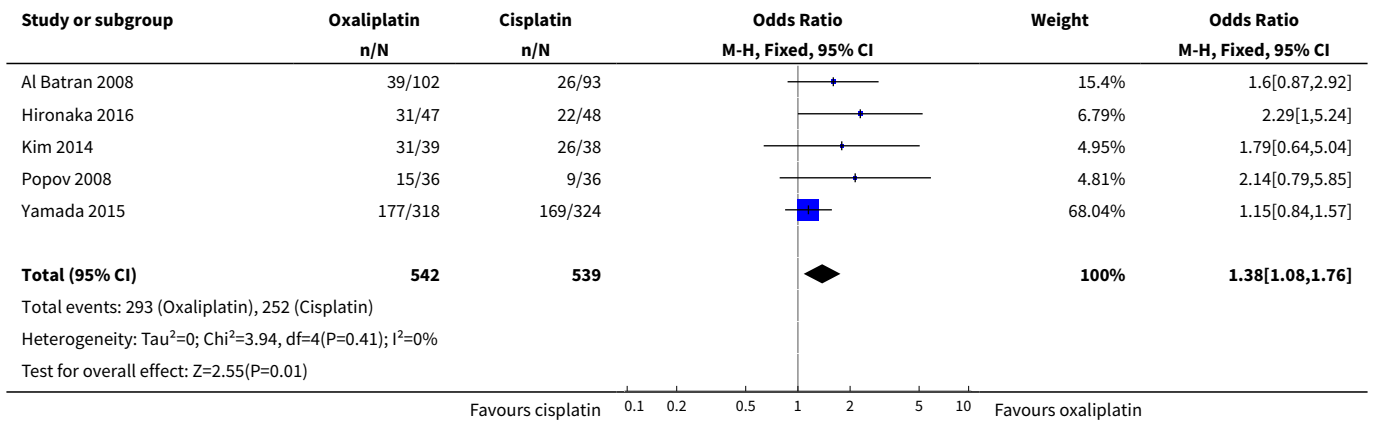
Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Overall Survival	5	1105	Hazard Ratio (Random, 95% CI)	0.81 [0.67, 0.98]
2 Tumour response	5	1081	Odds Ratio (M-H, Fixed, 95% CI)	1.38 [1.08, 1.76]
3 Progression-free survival	4	1034	Hazard Ratio (Random, 95% CI)	0.88 [0.66, 1.19]
4 Treatment-related death	5	1132	Odds Ratio (M-H, Fixed, 95% CI)	0.47 [0.17, 1.30]
5 Treatment discontinuation due to toxicity	3	970	Odds Ratio (M-H, Random, 95% CI)	0.97 [0.44, 2.13]

Analysis 8.1. Comparison 8 Chemotherapy with oxaliplatin versus the same regime including cisplatin, Outcome 1 Overall Survival.

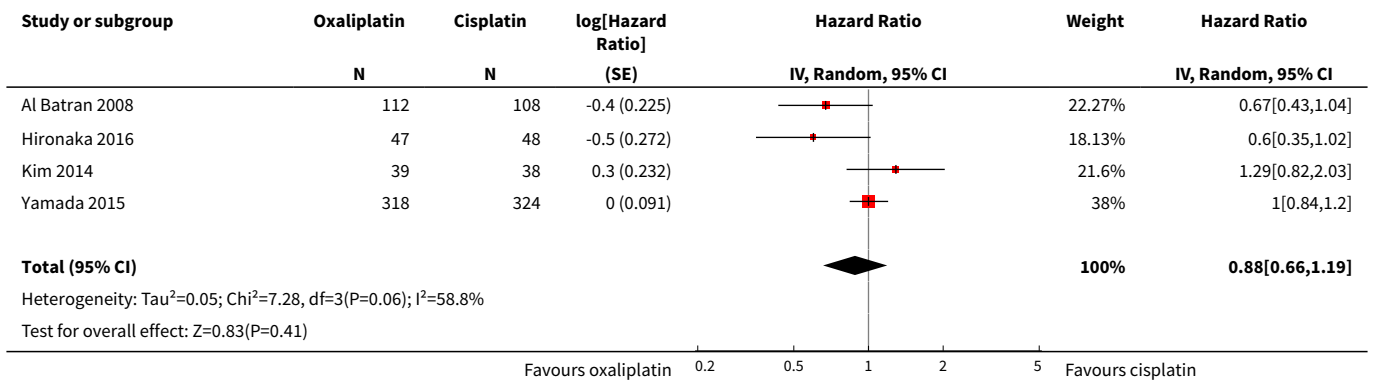




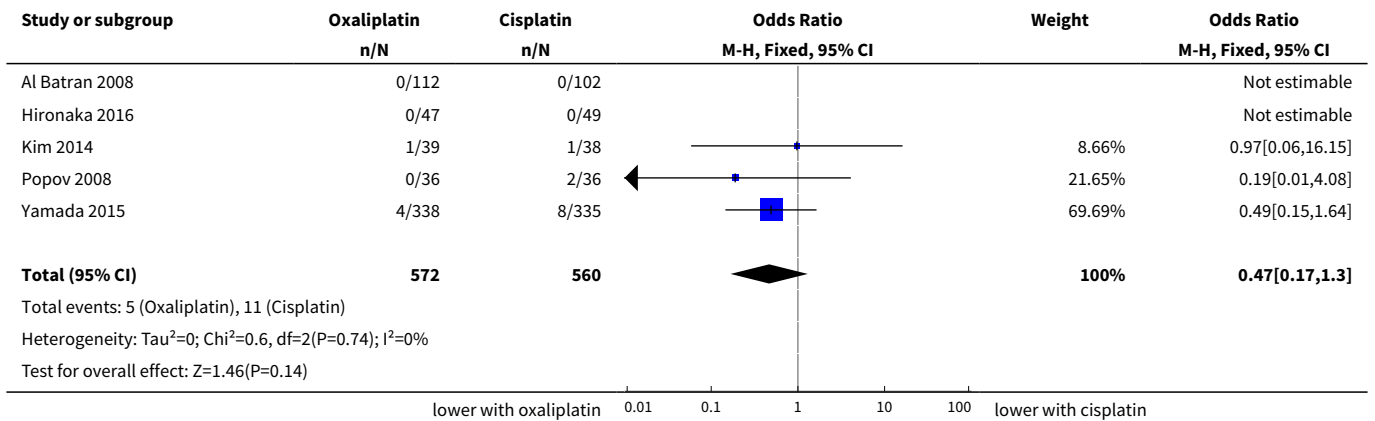
Analysis 8.2. Comparison 8 Chemotherapy with oxaliplatin versus the same regime including cisplatin, Outcome 2 Tumour response.



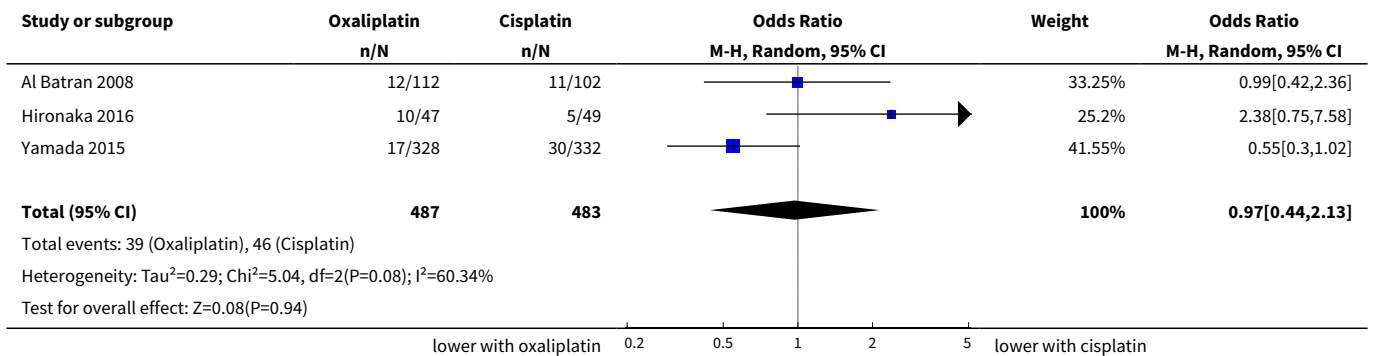
Analysis 8.3. Comparison 8 Chemotherapy with oxaliplatin versus the same regime including cisplatin, Outcome 3 Progression-free survival.



Analysis 8.4. Comparison 8 Chemotherapy with oxaliplatin versus the same regime including cisplatin, Outcome 4 Treatment-related death.



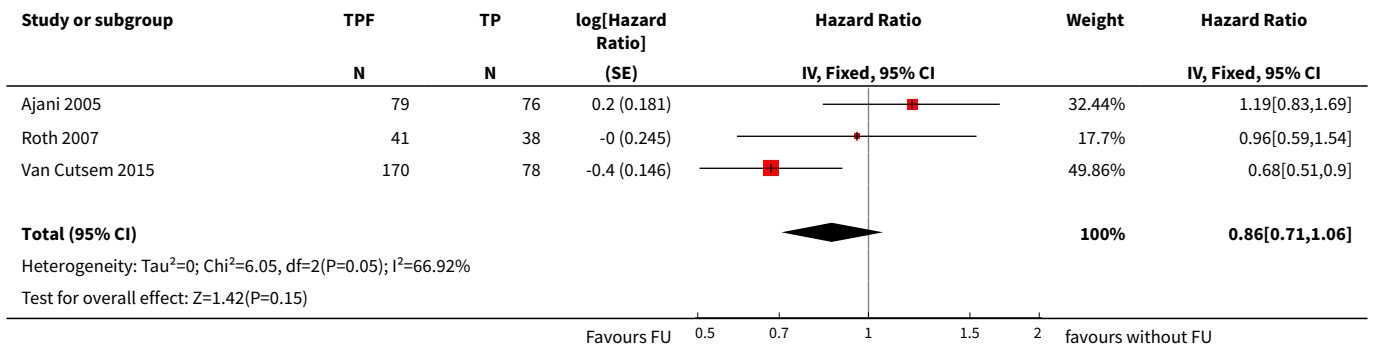
Analysis 8.5. Comparison 8 Chemotherapy with oxaliplatin versus the same regime including cisplatin, Outcome 5 Treatment discontinuation due to toxicity.



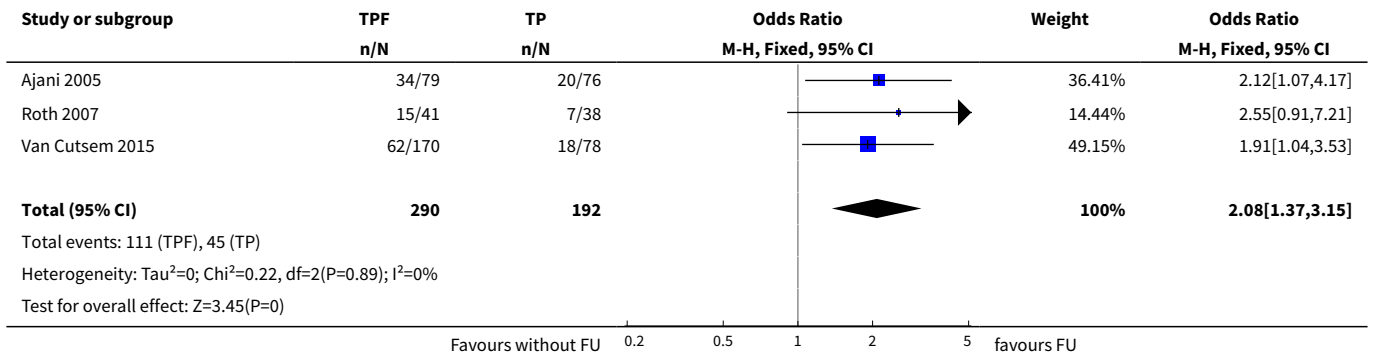
Comparison 9. Taxane-platinum-fluoropyrimidine combinations versus taxane-platinum (without fluoropyrimidine)

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Overall survival	3	482	Hazard Ratio (Fixed, 95% CI)	0.86 [0.71, 1.06]
2 Tumour response	3	482	Odds Ratio (M-H, Fixed, 95% CI)	2.08 [1.37, 3.15]
3 Progression-free survival	3	482	Hazard Ratio (Fixed, 95% CI)	0.74 [0.59, 0.93]
4 Treatment-related death	3	482	Odds Ratio (M-H, Fixed, 95% CI)	1.95 [0.73, 5.17]
5 Treatment discontinuation due to toxicity	2	234	Odds Ratio (M-H, Fixed, 95% CI)	1.71 [0.79, 3.69]

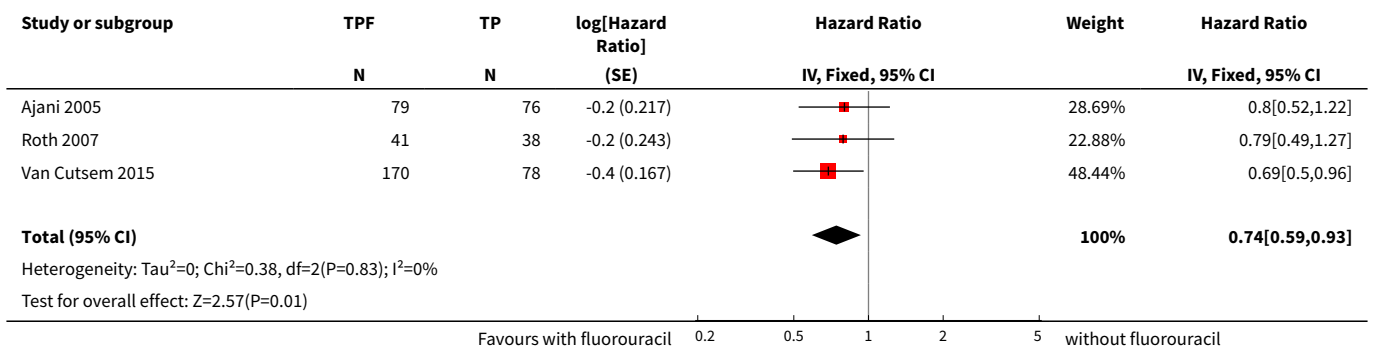
Analysis 9.1. Comparison 9 Taxane-platinum-fluoropyrimidine combinations versus taxane-platinum (without fluoropyrimidine), Outcome 1 Overall survival.



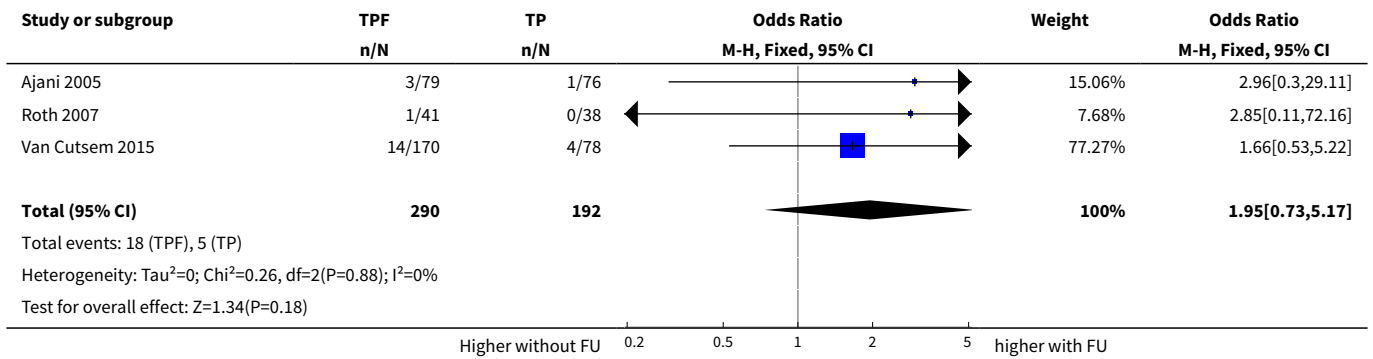
Analysis 9.2. Comparison 9 Taxane-platinum-fluoropyrimidine combinations versus taxane-platinum (without fluoropyrimidine), Outcome 2 Tumour response.



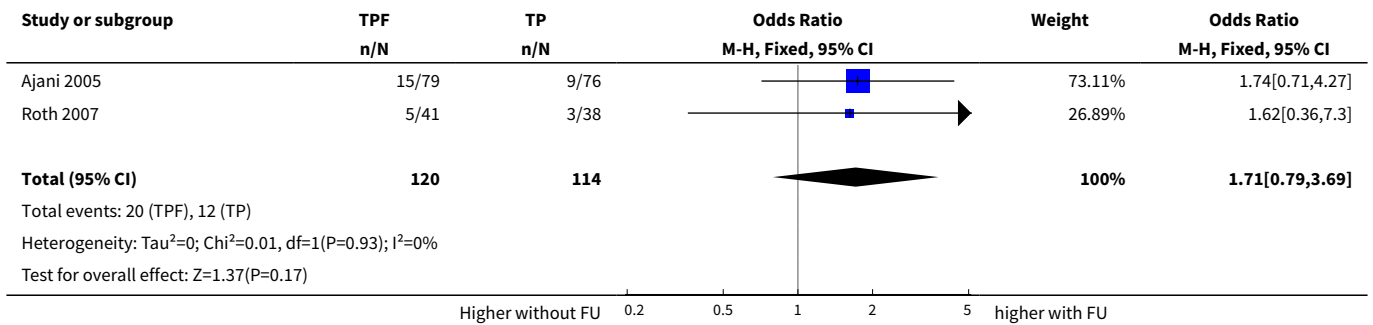
Analysis 9.3. Comparison 9 Taxane-platinum-fluoropyrimidine combinations versus taxane-platinum (without fluoropyrimidine), Outcome 3 Progression-free survival.



Analysis 9.4. Comparison 9 Taxane-platinum-fluoropyrimidine combinations versus taxane-platinum (without fluoropyrimidine), Outcome 4 Treatment-related death.



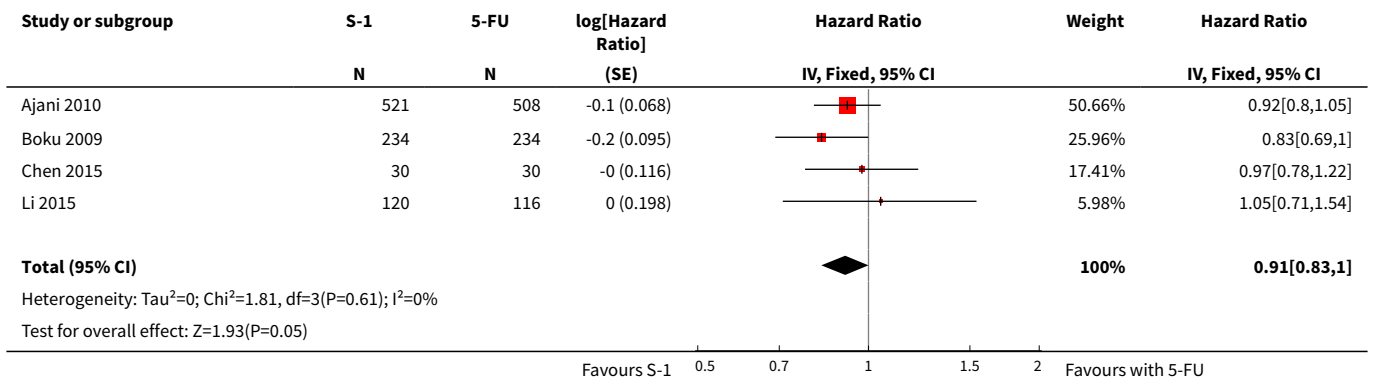
Analysis 9.5. Comparison 9 Taxane-platinum-fluoropyrimidine combinations versus taxane-platinum (without fluoropyrimidine), Outcome 5 Treatment discontinuation due to toxicity.



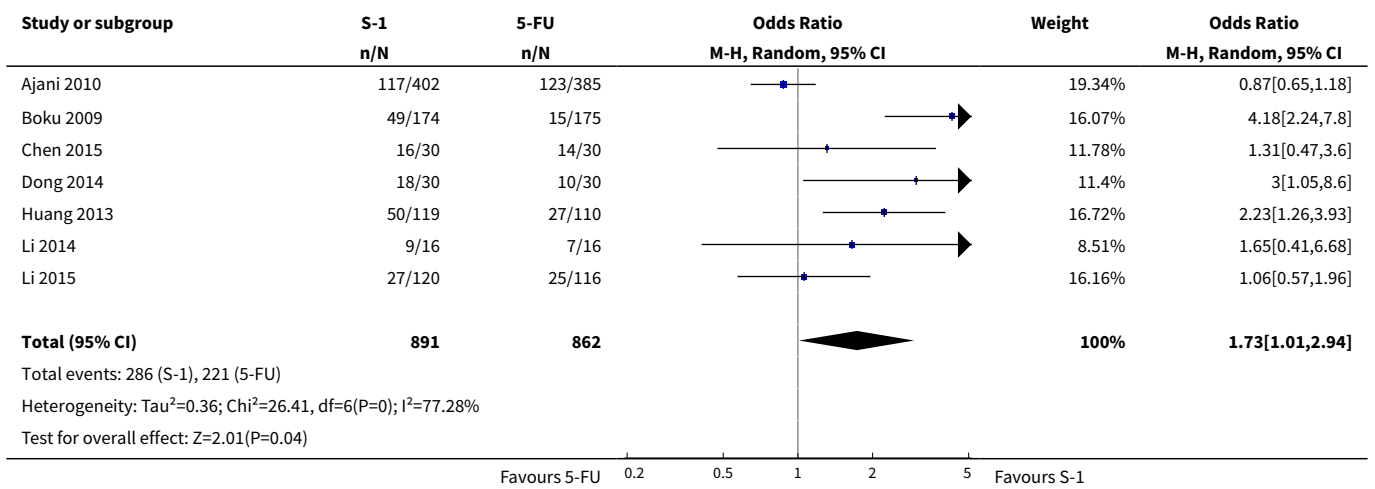
Comparison 10. S-1 versus 5-FU-containing regimes

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Overall Survival	4	1793	Hazard Ratio (Fixed, 95% CI)	0.91 [0.83, 1.00]
2 Tumour response	7	1753	Odds Ratio (M-H, Random, 95% CI)	1.73 [1.01, 2.94]
3 Progression-free survival	4	1942	Hazard Ratio (Random, 95% CI)	0.85 [0.70, 1.04]
4 Time-to treatment failure	5	1818	Hazard Ratio (Random, 95% CI)	0.88 [0.76, 1.01]
5 Treatment-related deaths	4	1962	Odds Ratio (M-H, Fixed, 95% CI)	0.56 [0.30, 1.06]
6 Treatment discontinuation due to toxicity	3	1726	Odds Ratio (M-H, Fixed, 95% CI)	0.85 [0.63, 1.13]

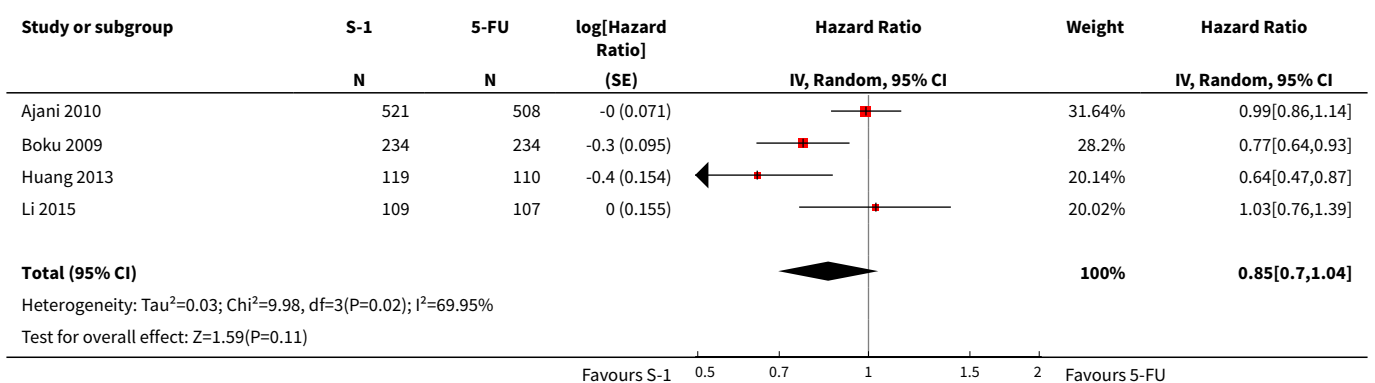
Analysis 10.1. Comparison 10 S-1 versus 5-FU-containing regimes, Outcome 1 Overall Survival.



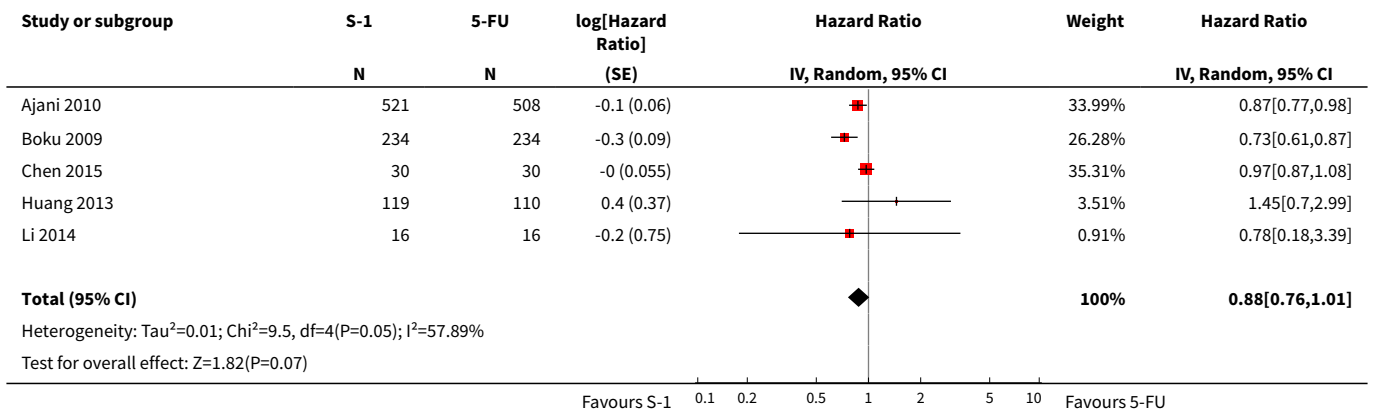
Analysis 10.2. Comparison 10 S-1 versus 5-FU-containing regimes, Outcome 2 Tumour response.



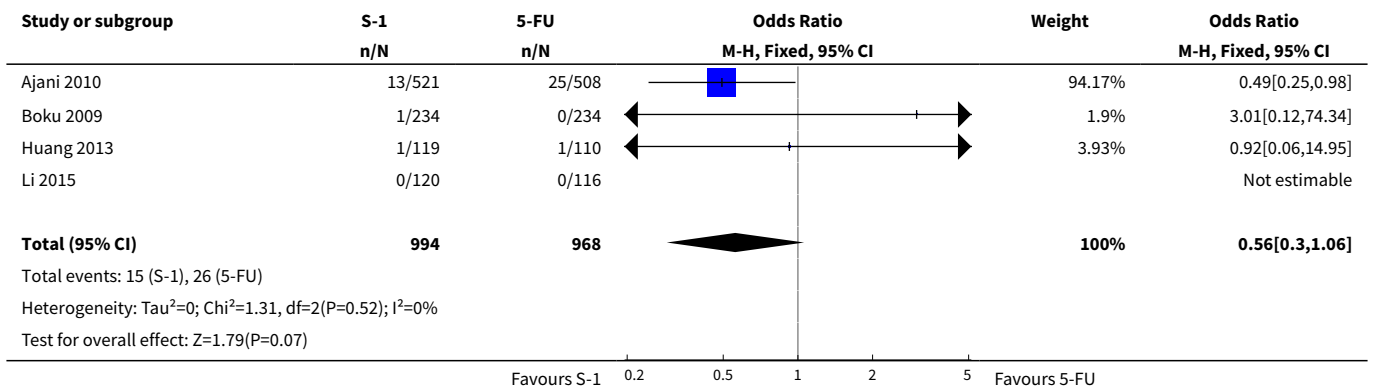
Analysis 10.3. Comparison 10 S-1 versus 5-FU-containing regimes, Outcome 3 Progression-free survival.



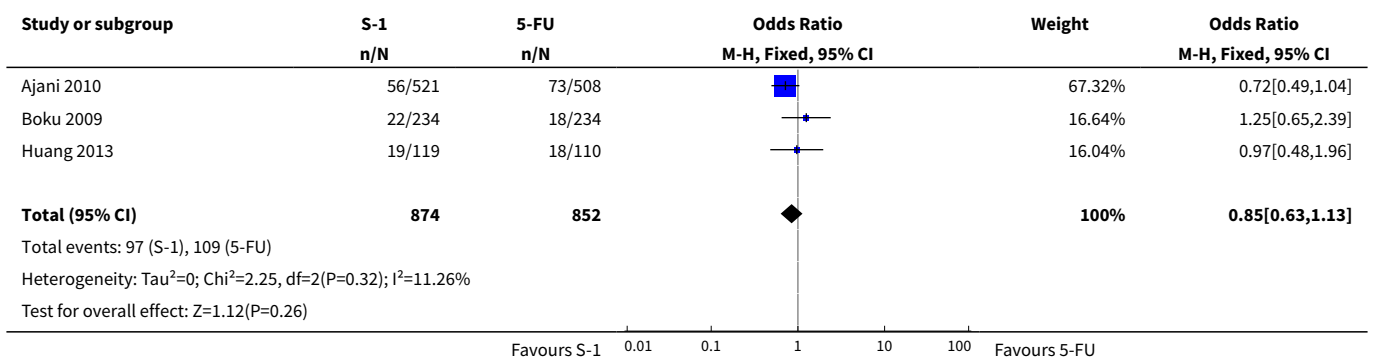
Analysis 10.4. Comparison 10 S-1 versus 5-FU-containing regimes, Outcome 4 Time-to treatment failure.



Analysis 10.5. Comparison 10 S-1 versus 5-FU-containing regimes, Outcome 5 Treatment-related deaths.



Analysis 10.6. Comparison 10 S-1 versus 5-FU-containing regimes, Outcome 6 Treatment discontinuation due to toxicity.



APPENDICES**Appendix 1. CENTRAL update January 2013**

1. exp stomach neoplasms/
2. (stomach adj5 neoplas\$).ti,ab.
3. (stomach adj5 cancer\$).ti,ab.
4. (stomach adj5 carcin\$).ti,ab.
5. (stomach adj5 tumo\$).ti,ab.
6. (stomach adj5 metasta\$).ti,ab.
7. (stomach adj5 malig\$).ti,ab.
8. (gastric adj5 neoplas\$).ti,ab.
9. (gastric adj5 cancer\$).ti,ab.
10. (gastric adj5 carcin\$).ti,ab.
11. (gastric adj5 tumo\$).ti,ab.
12. (gastric adj5 metasta\$).ti,ab.
13. (gastric adj5 malig\$).ti,ab.
14. or/1-13
15. exp drug therapy/
16. chemothera\$.ti,ab.
17. drug therap\$.ti,ab.
18. antineoplastic\$.ti,ab.
19. or/15-18
20. exp palliative care/
21. palliat\$.ti,ab.
22. unresect\$.ti,ab.
23. inopera\$.ti,ab.
24. advanc\$.ti,ab.
25. (best adj5 support\$ adj5 care).ti,ab.
26. unopera\$.ti,ab.
27. (non adj5 resect\$).ti,ab.
28. nonresect\$.ti,ab.
29. or/20-28
30. 14 and 19
31. 29 and 30
32. limit 31 to yr="2009 - 2013"

Appendix 2. MEDLINE update March 2009-Jan 2013

1. randomized controlled trial.pt.
2. controlled clinical trial.pt.
3. randomized.ab.
4. placebo.ab.
5. drug therapy.fs.
6. randomly.ab.
7. trial.ab.
8. groups.ab.
9. or/1-8
10. exp animals/ not humans.sh.
11. 9 not 10
12. exp stomach neoplasms/
13. (stomach adj5 neoplas\$).tw.
14. (stomach adj5 cancer\$).tw.
15. (stomach adj5 carcin\$).tw.
16. (stomach adj5 tumo\$).tw.
17. (stomach adj5 metasta\$).tw.
18. (stomach adj5 malig\$).tw.
19. (gastric adj5 neoplas\$).tw.
20. (gastric adj5 cancer\$).tw.
21. (gastric adj5 carcin\$).tw.
22. (gastric adj5 tumo\$).tw.
23. (gastric adj5 metasta\$).tw.
24. (gastric adj5 malig\$).tw.
25. or/12-24
26. exp drug therapy/
27. chemothera\$.tw.
28. drug therap\$.tw.
29. antineoplastic\$.tw.
30. or/26-29
31. exp palliative care/
32. palliat\$.tw.
33. unresect\$.tw.
34. inopera\$.tw.

35. advanc\$.tw.
36. (best adj5 support\$ adj5 care).tw.
37. unopera\$.tw.
38. (non adj5 resect\$.tw.
39. nonresect\$.tw.
40. or/31-39
41. 25 and 30
42. 40 and 41
43. 42 and 11
44. limit 43 to ed=20090309-20130131

Appendix 3. Embase update March 2009-Jan 2013

1. exp randomized controlled trial/
2. randomi?ed controlled trial\$.tw.
3. exp randomization/
4. exp single blind procedure/
5. exp double blind procedure/
6. or/1-5
7. animal.hw.
8. human.hw.
9. 7 not (7 and 8)
10. 6 not 9
11. exp clinical trial/
12. (clin\$ adj3 stud\$).ti,ab,tw.
13. (clin\$ adj3 trial\$).ti,ab,tw.
14. ((singl\$ or doubl\$ or treb\$ or tripl\$) adj3 (blind\$ or mask\$)).ti,ab,tw.
15. exp placebo/
16. placebo\$.ti,ab,tw.
17. random.ti,ab,tw.
18. (crossover\$ or cross-over\$).ti,ab,tw.
19. or/11-18
20. 19 not 9
21. 20 not 10
22. exp comparative study/
23. exp evaluation/
24. exp prospective study/

25. exp controlled study/
26. (control\$ or prospective\$ or volunteer\$).ti,ab,tw.
27. or/22-26
28. 27 not 9
29. 10 or 21 or 28
30. exp stomach tumor/
31. (stomach adj5 neoplas\$).tw.
32. (stomach adj5 cancer\$).tw.
33. (stomach adj5 carcin\$).tw.
34. (stomach adj5 tumo\$).tw.
35. (stomach adj5 metasta\$).tw.
36. (stomach adj5 malig\$).tw.
37. (gastric adj5 neoplas\$).tw.
38. (gastric adj5 cancer\$).tw.
39. (gastric adj5 carcin\$).tw.
40. (gastric adj5 tumo\$).tw.
41. (gastric adj5 metasta\$).tw.
42. (gastric adj5 malig\$).tw.
43. or/30-42
44. exp drug therapy/
45. chemothera\$.tw.
46. drug therap\$.tw.
47. antineoplastic\$.tw.
48. or/44-47
49. exp palliative therapy/
50. palliat\$.tw.
51. unresect\$.tw.
52. inopera\$.tw.
53. advanc\$.tw.
54. (best adj5 support\$ adj5 care).tw.
55. unopera\$.tw.
56. (non adj5 resect\$).tw.
57. nonresect\$.tw.
58. or/49-57
59. 43 and 48

60. 58 and 59

61. 29 and 60

62. limit 61 to em=200910-201306

WHAT'S NEW

Date	Event	Description
30 June 2016	New search has been performed	Searches rerun and results incorporated
30 June 2016	New citation required and conclusions have changed	We added 26 included studies, two new comparisons (9) Taxane-platinum-fluoropyrimidine combinations versus taxane-platinum (without fluoropyrimidine) and (10) S-1 versus 5-FU containing regimens and subgroup analyses to comparisons 5 and 6; Irinotecan versus non-irinotecan-containing regimens and docetaxel versus non-docetaxel-containing regimens

HISTORY

Protocol first published: Issue 1, 2003

Review first published: Issue 2, 2005

Date	Event	Description
6 December 2009	New citation required and conclusions have changed	Conclusions changed, authors changed.
3 December 2009	New search has been performed	Updated, new studies added and change of statistical model due to heterogeneity.
30 October 2008	Amended	Converted to new review format.
28 May 2004	Amended	Conclusions changed.
1 February 2004	Amended	New studies found and included or excluded.

CONTRIBUTIONS OF AUTHORS

Wagner AD: Protocol development, eligibility and quality assessment, data extraction and analysis, drafting of final review.

Unverzagt S: Eligibility and quality assessment in the update, responsible statistician of the current update. Data extraction and analysis, and drafting of the final review (2017 update).

Moehler, M: Review of search results, drafting of the final review (2017 update).

Grothe W: Literature searches, eligibility and quality assessment, data extraction and analysis.

Haerting J: Protocol development, statistician in initial version of the review.

Syn, NL: Review of search results, quality assessment, data extraction and analysis, drafting of final review (2017 update).

Ho, J: Review of search results, data extraction and analysis, drafting of final review (2017 update).

Yong, WP: Review of search results, drafting of final review (2017 update).

Tai, BC: Data extraction and analysis (2017 update).

DECLARATIONS OF INTEREST

ADW: has received consulting fees from Eli Lilly, Celgene, MerckKG/Phizer, MSD, TAIHO and Roche pharmaceuticals for consultancy work over the past three years, travel and accomodation support from Janssen and support from Merck Serono to conduct a clinical trial of

targeted treatment for gastric cancer, which is beyond the scope of this review. The overall number of consultancy agreements per year is low (usually less than five advisory boards/year, with fees of between 500 and 1500 Euro per event). All funds were paid to her institution in accordance with their guidelines. A portion of the money that the institution receives is transferred to ADW and represents a trivial percentage of her salary.

NLXS: none known.

MM: has received consulting fees from Eli Lilly, Onyx, Roche, Nordic, Amgen, MSD, Merck Serono, Pfizer and BMS. He has received payment for lectures from Falk, Nordic, Amgen, mci, MSD, Merck Serono, Pfizer and BMS. His institution has received funds for independent research grants from Merck, Amgen, BMS, Taiho, Roche, AIO, MSD, and the EORTC.

WG: none known.

WPY: has received payment from Eli Lilly for service on speaker bureaus and travelling/accommodation for conferences from BMS and Taiho, unrelated to this review.

BCT: has received a payment from Boehringer Ingelheim for providing a lecture, and royalties from Wiley-Blackwell from a publication. Her institution has received funds from the Health Science Authority (Singapore) for statistical consultancy.

JH: none known.

SU: none known.

SOURCES OF SUPPORT

Internal sources

- Department of Internal Medicine I, Martin-Luther-University Halle-Wittenberg, Germany.
- Department of Internal Medicine IV, Martin-Luther-University Halle-Wittenberg, Germany.
- Institute of Medical Epidemiology, Biometry and Informatics, Martin-Luther-University, Halle-Wittenberg, Germany.
- Co-ordinating Centre for Clinical Trials, Halle (supported by German Ministry of Education & Research, Grant Number BMBF/FKZ: 01GH0105 KKS, Halle), Germany.

External sources

- No sources of support supplied

DIFFERENCES BETWEEN PROTOCOL AND REVIEW

There were a large number of categories of different combination chemotherapy regimens and the number of relevant studies in each category was not known when writing the protocol. For this reason we were unable to plan in advance the best way to compare categories of all relevant categories of combination chemotherapies (Van Cutsem 2009).

We clarified how we dealt with multiple study arms from one study.

In the first update of this review (Wagner 2010, submitted in November 2009), we chose to add the following comparisons based on their clinical relevance and the availability of a sufficient number of relevant studies. Among these, comparisons 3 and 4 have been added in the discussion of the previously published version of this review as they were not specified in advance. Comparisons 5 to 8 have been specified in advance in the last update of the review.

In this second update of this review (submitted in March 2017), we added two additional comparisons (9 and 10) and we differentiated between additive and substitutive comparisons in comparisons 5 and 6.

Comparison number:

3. 5-FU/cisplatin/anthracycline combinations versus 5-FU/cisplatin (without anthracyclines)
4. 5-FU/cisplatin/anthracycline-combinations versus 5-FU/anthracycline combinations (without cisplatin)
5. Irinotecan versus non-irinotecan-containing regimens
6. Docetaxel versus non-docetaxel-containing regimens
7. Regimens including oral 5-FU prodrugs versus intravenous fluoropyrimidines
8. Oxaliplatin versus cisplatin-containing regimens
9. Taxane-platinum-fluoropyrimidin combinations versus taxane-platinum (without fluoropyrimidine)
10. S-1 versus 5-FU-containing regimens

The method for statistical analysis in case of significant heterogeneity ($I^2 > 20\%$) is a random-effects model. Although the original version of the protocol does not state this, these changes were requested after methodological review by the Cochrane UGPD group. This methodological change is responsible for differences in HRs between this and the previously published version of the review (Wagner 2006).

As recent studies have increasingly reported progression-free survival instead of time to progression, we included the former as secondary endpoints in this update.

While there was no evidence for both - the use of second-line therapy and targeted therapies - at the time the protocol for this review was written, both second-line and these targeted agents (trastuzumab in first-line for HER-2 positive people and ramucirumab in second-line) have changed the management options for advanced gastric cancer significantly and are therefore included in the discussion of the treatment options although not analysed in detail by this review.

In this updated review, we no longer use the Jadad scale to assess methodological quality (Jadad 1996). Instead we independently assessed the risk of bias in the included studies using the 'Risk of bias' assessment tool described in Chapter 8 of the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011).

INDEX TERMS

Medical Subject Headings (MeSH)

Anthracyclines [administration & dosage]; Antineoplastic Combined Chemotherapy Protocols [*therapeutic use]; Camptothecin [administration & dosage] [analogs & derivatives]; Cisplatin [administration & dosage]; Docetaxel; Fluorouracil [administration & dosage]; Irinotecan; Randomized Controlled Trials as Topic; Stomach Neoplasms [*drug therapy] [mortality]; Taxoids [administration & dosage]

MeSH check words

Humans