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# The effect of pre-anaesthetic assessment clinic: a systematic review of randomised and non-randomised prospective controlled studies

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3 4	1	The effect of pre-anaesthetic assessment clinic: a systematic review of randomised and						
5 6 7	2	non-randomised prospective controlled studies						
8 9	3							
10 11 12	4	Eirunn Wallevik Kristoffersen <sup>1,</sup> *, Anne Opsal <sup>1</sup> , Tor Tveit <sup>1,2,3</sup> , Rigmor C Berg <sup>4,5</sup> , Mariann						
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2 3	24	ABSTRACT					
4 5	27	<b>Objectives:</b> The sim of this systematic review was to examine the effectiveness of pro					
6	25	apportation association of this systematic review was to examine the effectiveness of pre-					
7 8	20	anaesthetic assessment clinics (PACs) implemented to improve quality and patient safety in					
9 10	27	perioperative care.					
11 12 13	28	Design: Systematic review.					
	29	Data sources: The electronic databases CINAHL Plus with Full Text (EBSCOhost), Medline,					
14 15	30	and Embase (OvidSP) were systematically searched from 1st April, 1996 to 4th February,					
16 17	31	2021.					
18	32	Eligibility criteria: The main inclusion criterion was that the study, using empirical					
19 20	33	quantitative methods, addressed the effectiveness of PACs.					
21 22	34	Data extraction and synthesis: Titles, abstracts, and full texts were screened in duplicate by					
23 24	35	two authors. Risk of bias assessment, using the Joanna Briggs Institute critical appraisal					
25	36	checklist for quasi-experimental studies, and data extraction were performed by one author					
20	37	and checked by the other author. Results were synthesised narratively owing to the					
28 29	38	heterogeneity of the included studies.					
30 31	39	Results: Seven prospective controlled studies were conducted. Most studies had a high risk					
32 33	40	of bias. Three studies reported a significant reduction in the length of the hospital stay, and					
34 35	41	two studies reported a significant reduction in cancellation of surgery for medical reasons					
36 37	42	when patients were seen in the PAC. In addition, the included studies presented mixed					
38	43	results regarding anxiety in patients.					
39 40	44	Conclusion: This systematic review demonstrated a reduction in the length of hospital stay					
41 42	45	and cancellation of surgery when the patients had been assessed in the PAC. There is a need					
43 44	46	for high-quality prospective studies to gain a deeper understanding of the effectiveness of					
45 46	47	PACs.					
40	48	PROSPERO registration number: CRD42019137724					
48 49	49						
50 51	50	Keywords: pre-anaesthetic assessment clinic, preoperative care, quality, safety, systematic					
52 53	51	review					
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4 5	59	ARTICLE SUMMARY
6	60	
7	61	Strengths and Limitations of this study
o 9	62	Only prospective studies were included in this systematic review.
10 11	63	• The systematic review was conducted in accordance with international guidelines.
12 13	64	• Only seven studies were identified, highlighting the need for further research on pre-
14 15	65	anaesthetic assessment clinics.
16 17	66	<ul> <li>Overall, the quality of the included studies was low, and the current practice</li> </ul>
18 19	67	possesses limited evidence base.
20 21	68	
22	69	INTRODUCTION
24 25 26	70	Anaesthesia constitutes an important part of surgery; however, it has the potential to
20 27 28	71	activate physiological changes that can increase morbidity and mortality,[1] mainly
29 30	72	depending on the patients' preoperative health condition and age.[2] Hospitals are treating
31 32 33	73	patients with complex, comorbid healthcare problems who undergo progressively extensive
34 35	74	surgeries and interventions.[3,4] To ensure the quality and safety of anaesthesia and
36 37 38	75	surgery, precise knowledge of the clinical characteristics of patients undergoing surgery is
39 40	76	critical to the perioperative treatment plan.[2] Over the past 50 years, perioperative
41 42 43	77	mortality, including anaesthesia-related mortality, has declined, with the most significant
44 45	78	decline observed in developed countries, [1,5] mainly due to new anaesthetics, improved
46 47 48	79	monitoring equipment and training, availability of recovery rooms, and improved airway
49 50	80	management.[4] However, an Australian study reported that 14% of anaesthetic-surgical
51 52 53	81	complications and 39% of deaths attributed to anaesthesia were associated with insufficient
55 55	82	and/or inadequate preoperative evaluation.[6] A Danish retrospective investigation showed
56 57	83	that the deaths among patients undergoing surgery could have been prevented by a
58 59 60	84	thorough preoperative evaluation,[7] indicating that risk factors are both patient-and

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85	surgery-related and linked to organisational structures.[8] Future efforts should improve
86	preoperative anaesthesia safety, [9] by improving planning and preparation for elective
87	procedures and interventions.

88 In 1949, Lee discussed the value of the "anaesthetic outpatient clinic" in the preparation of 89 patients for surgery.[10] Today, an increasing number of pre-anaesthesia assessment clinics 90 (PACs) are supporting hospitals in handling the rise in the number and complexity of surgical 91 procedures.[11] The PAC consultation, conducted by the anaesthesiologist, anaesthesia 92 nurse, or both, is globally recognised as an evaluation method while optimising the patients' 93 medical condition prior to surgery and anaesthesia, and is considered essential in securing 94 anaesthetic practice since it detects anaesthesia-related risk factors and high-risk patients, 95 improves patient outcomes, prepares the patient physically and psychologically for 96 anaesthesia, and ensures the patient's most favourable condition for surgery and 97 anaesthesia.[12-14] Considering the well-prepared patients and staff, several researchers 98 posit that with PAC, the number of surgical cancellations, length of hospital stay, and 99 mortality rate are reduced, and tests are minimised. [8,15,16] Others assert that patients feel 100 less anxious regarding the subsequent anaesthetic and surgical processes and are highly 101 satisfied with this service when PACs are used. [15, 17, 18]

As Turunen *et al.* state, research on PACs is scarce regarding costs, financial savings, the impact on patient safety and quality of care, accuracy of operative patients, and effect on preoperative nursing levels.[19] Survey results indicate that anaesthesiologists perceive day of surgery delays due to missing information as common, even with PAC consultations.[20] The present systematic review examines the outcomes of PAC as systematic work on quality BMJ Open

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3 4	107	and patient safety, including identifying the areas for improvement, implementing							
5 6 7 8	108	interventions, and ensuring that patient outcome improvement.							
9 10	109	METHODS							
11 12 13	110	The aim of this systematic review was to examine the effectiveness of PACs in improving							
14 15 16	111	quality and patient safety in preoperative care. A further aim was to determine the gaps in							
16 17 18	112	existing knowledge for future research. Our systematic review followed the guidelines in the							
19 20 21	113	Cochrane Handbook for Systematic Reviews of Interventions [21] and was reported in							
21 22 23	114	accordance with the Preferred Reporting Items for Systematic Reviews and Meta-Analyses							
24 25 26	115	(PRISMA) guidelines.[22] The protocol was published in PROSPERO: CRD42019137724.[23]							
20 27 28	116	We had two review questions:							
29 30 31	117	1. What are the effects of PACs on patient satisfaction, anxiety, and safety?							
32 33 34 25	118	2. What are the effects of PACs on cancellation rate, cost, and efficiency?							
35 36 37	119	Search strategies							
38 39 40	120	We performed a scoping search in different databases to identify the key terms for the							
41 42	121	literature search.[24,25] The final search was planned and conducted in close collaboration							
43 44 45	122	with a university librarian. On 11th September, 2018 we searched CINAHL Plus with Full Text							
46 47 48	123	(EBSCOhost), Medline, and Embase (OvidSP), and updated it on 4th February, 2021.							
49 50	124	Considering the lack of subject headings (e.g., MeSH) for PAC, we used text words such as							
51 52 53	125	preanaesthesia. The search in Medline is presented in Appendix 1. The search mode in							
54 55	120	cinant was boolean/Phase, which supports boolean searching or exact phrase searching. To							
56 57 58	127	limited the search to 1996 since this was the year one of the first known articles in this area							
59 60	120	innited the search to 1990 since this was the year one of the first known articles in this died							

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3 4	129	was published.[23] Complementary methods to identify studies included following up on
5 6 7	130	citations via Scopus, scanning the reference lists of relevant papers and included articles,
7 8 9	131	and checking for relevant studies in clinical trials.[24]
10 11 12	132	Eligibility criteria
13 14	133	Considering the aim of the review, the main inclusion criterion was that the study, using
15 16 17	134	empirical quantitative methods, addressed the effectiveness of PACs. Specific study
17 18 19	135	eligibility criteria were: (a) published in English or Scandinavian language, (b) scientific
20 21	136	publication of original research, (c) reporting the outcomes of PAC, (d) PAC consultation with
22 23 24	137	the patient present, (e) randomised or non-randomised prospective controlled studies, and
25 26	138	(f) newly established PAC. We excluded: (a) editorials, discussion papers, and conference
27 28 20	139	abstracts, (b) reviews, (c) instrument testing, (d) studies with children, and (e) retrospective
29 30 31 32 33	140	studies.
33	141	Study selection
33 34 35 36	141 142	Study selection All references identified in the search were transferred to EndNoteX9, where the duplicates
32 33 34 35 36 37 38	141 142 143	Study selection All references identified in the search were transferred to EndNoteX9, where the duplicates were removed. Next, all unique references were transferred to the Covidence screening
32 33 34 35 36 37 38 39 40 41	141 142 143 144	Study selection         All references identified in the search were transferred to EndNoteX9, where the duplicates         were removed. Next, all unique references were transferred to the Covidence screening         tool.[26] Study eligibility was ascertained independently by two authors, first at the title and
32 33 34 35 36 37 38 39 40 41 42 43	141 142 143 144 145	Study selection All references identified in the search were transferred to EndNoteX9, where the duplicates were removed. Next, all unique references were transferred to the Covidence screening tool.[26] Study eligibility was ascertained independently by two authors, first at the title and abstract level, and subsequently at the full text. Inclusion was determined by consensus, and
32 33 34 35 36 37 38 39 40 41 42 43 44 45 46	141 142 143 144 145 146	Study selection All references identified in the search were transferred to EndNoteX9, where the duplicates were removed. Next, all unique references were transferred to the Covidence screening tool.[26] Study eligibility was ascertained independently by two authors, first at the title and abstract level, and subsequently at the full text. Inclusion was determined by consensus, and disagreements were resolved by consulting a third author.
32 33 34 35 36 37 38 39 40 41 42 43 44 45 46 47 48	141 142 143 144 145 146 147	Study selection All references identified in the search were transferred to EndNoteX9, where the duplicates were removed. Next, all unique references were transferred to the Covidence screening tool.[26] Study eligibility was ascertained independently by two authors, first at the title and abstract level, and subsequently at the full text. Inclusion was determined by consensus, and disagreements were resolved by consulting a third author. Quality assessment
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32         33         34         35         36         37         38         39         40         41         42         43         44         45         46         47         48         49         50         51         52         53         54         55	141 142 143 144 145 146 147 148 149 150	Study selection All references identified in the search were transferred to EndNoteX9, where the duplicates were removed. Next, all unique references were transferred to the Covidence screening tool.[26] Study eligibility was ascertained independently by two authors, first at the title and abstract level, and subsequently at the full text. Inclusion was determined by consensus, and disagreements were resolved by consulting a third author. Quality assessment We used design-specific checklists to assess the studies' risk of bias. Given the methodological similarity of the included studies, only the Joanna Briggs Institute Critical appraisal checklist for quasi-experimental studies was used.[27] One author performed the
32         33         34         35         36         37         38         39         40         42         43         44         45         46         47         48         90         51         52         54         55         56         57         58	141 142 143 144 145 146 147 148 149 150 151	Study selection All references identified in the search were transferred to EndNoteX9, where the duplicates were removed. Next, all unique references were transferred to the Covidence screening tool.[26] Study eligibility was ascertained independently by two authors, first at the title and abstract level, and subsequently at the full text. Inclusion was determined by consensus, and disagreements were resolved by consulting a third author. Quality assessment We used design-specific checklists to assess the studies' risk of bias. Given the methodological similarity of the included studies, only the Joanna Briggs Institute Critical appraisal checklist for quasi-experimental studies was used.[27] One author performed the risk of bias assessment, and the other checked the accuracy of the assessment.

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Disagreements were resolved through discussion with a third author. Each of the nine checklist questions was answered no, yes, unclear (or not applicable).

### Data extraction and analysis

One author extracted data from each included study onto a pre-designed Excel spreadsheet, and another checked the extracted data for accuracy, consistency, and completeness. Extracted information included publication details, study design, setting, and characteristics of the patients, interventions, comparisons, and outcome (PICO). We requested information on the missing data; however, received no response from the author. If the PICO elements were sufficiently similar and statistical data were available, we had planned to conduct meta-analyses. However, the extracted data revealed substantial heterogeneity among the studies, and there were no randomised controlled trials (RCTs). Therefore, we performed a narrative synthesis, describing and comparing the main findings from the included studies, and discussing their methodological strengths and weaknesses.

RESULTS

Figure 1. provides details of the study selection process. A total of 2250 records were dentified in the first search and 742 in the second search. After removing duplicates, we screened 2372 records based on the title and abstract; of these, 179 records passed the fulltext screening. We included seven studies that met the inclusion criteria.

## **Overall characteristics of the studies**

The seven included studies are listed in Table 1. They were all in English and published in 2000–2017, with data collected in the years 1997–2015 (one did not provide this data collection information).[28] Based on our inclusion criteria, all were prospective controlled studies, but we found no RCTs. There was one controlled before-after study.[34] The other six studies had control groups but no baseline assessments, only assessments following PAC

77411 patients.

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implementation. There were three 2-group non-parallel after-only studies, [29, 30, 32] and

three 2-group parallel after-only studies [28], where one had a matched control group[31]

and one had three follow-up assessments of one arm.[33] In total, the studies included

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## **Table 1: Description of included studies**

Author, Year, Country	Study design	Sampling time	Population	Intervention	Comparison	Outcomes
Farasatkish, 2009[1] Iran	2-group after study	May 2007 through August 2007	N=1716, open- heart surgery, ASA class III-IV	Pre-anaesthesia consultation clinic (3-10 days before surgery)	Usual care (within 24 h of surgery)	Cancellations
Kamal, 2011[2] England	2-group after study	April 2005 through April 2009	N=1445, complex elective orthopaedic surgery, ASA class III-IV	Preoperative anaesthetic assessment clinic (timing not stated)	Usual care (day of surgery)	Admissions, length of stay, mortality, cost
Kamau, 2017[3] Kenya	СВА	August 2000, April 2001, November 2001	N=51, elective non-cardiac surgery, ASA class III	Pre-anaesthesia clinic consultation (≥48 h before surgery)	Usual care (day before surgery)	Anxiety (STAI score)
Klopfenstein, 2000[4] Switzerland	2-group after study (parallel)	No data	N=40, elective endoscopic urological surgery, ASA class I-III	Pre-anaesthetic consultation (1-2 weeks before surgery)	Usual care (the evening before surgery)	Anxiety (MAACL, VAS)
Lee, 2012[5] China	2-group after study (parallel)	March 2007 through November 2009	N=352, elective surgery, ASA class I-IV	Anaesthesia consultation clinic (≤3 months before surgery)	Usual care (the evening before surgery)	Quality of recovery score), cost, cancellations, length of stay, satisfaction, anxiety (VAS), willingness to pay (WTP)

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3		Mendes,	2-group	April 2007	N=52254,	Preoperative	Usual care (timing	Cancellations,
4		2005[6]	after study	through	surgery, ASA	outpatient evaluation	not stated)	length of stay
5		Brazil	(parallel)	June 2007	class not stated	clinic (timing not	,	
7			(1			stated)		
8		van Klei	2-group	November	N=21553	Preoperative	Usual care (day	Cancellations same-
9		2002[7]	2 Broup	2012	elective surgery	outpatient evaluation	before surgery)	day admissions
10		2002[7] Tho	arter study	2012	ASA class mainly	clinic (avorage 2 wooks	before surgery	longth of stay
11		Nothorlands				before surgery)		length of stay
12	101	Nethenanus			1-11	before surgery)		
13 14	181							
15								
16	182	ASA: American S	ociety of Anesth	iesiology; CBA: co	ontrolled before-after	; MAACL: Multiple Affect A	Adjective Check List; ST	Al: State-Trait Anxiety
17	183	Inventory; VAS:	visual analogue s	scale;				
18	184	WTP: willingness	s to pay					
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Considering the intervention, PACs in all studies consisted of an outpatient service whereby patients were checked for medical conditions that are important for anaesthesia and informed regarding what to expect on the day of surgery. However, the terminology used for PACs varied; they served different surgical specialities, and the pre-anaesthesia consultation was conducted from ≥48 h to ≤3 months before the surgery. Three were implemented in a university hospital,[31,33,34] one in a teaching hospital,[30] one in a medical centre,[32] and one in a general hospital[29] (one study did not specify the context).[28] The person conducting the pre-anaesthesia consultation also varied: in five studies, it was the anaesthesiologists,[28-31,33] in the other studies it was (also) the orthopaedic senior house officer,[29] the consultant or resident,[34] or the physician.[32] In three studies, nurses were part of the team assessing the patients.[29-31] The comparison group in all studies was usual care, which generally involved performing a preoperative anaesthetic evaluation the day before the surgery on the admitted patients.

Of the 77411 patients in the studies, 9626 and 15531 patients were in the intervention and control groups, respectively. One study did not specify the number of patients in the intervention and control groups, but only the total number of surgeries performed.[33] Five studies reported data for sex, showing that 51% of the patients were women and 49% were men (12129 vs. 11583).[28,30-32,34] There were more females than males both in the intervention (4345 vs. 4134) and the control groups (7784 vs. 7449). Five studies reported data for age showing that all the patients were over 20 years old and grouped within the American Society of Anesthesiology (ASA) category.[28,30-32,34]

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The patients were scheduled to undergo a variety of surgeries, including orthopaedic,[29-31,34] urology,[28,30,31,34] general,[30,31,34] heart,[32] gynaecology/obstetrics, [30,31,34] vascular surgery,[30] ophthalmology,[30] maxillofacial/dental surgery, [30,34]neurological surgery,[30] and one did not specify the type of surgery.[33] In five studies, the type of anaesthesia was not specified,[29,30,32-34] and two studies reported patients for general and/or regional supplement.[28,31]

The patients included had previous anaesthetic experience in one study,[28] previous and no previous anaesthetic experience in another,[34] and five studies did not report this data.[29-33] Limited background characteristics of the patients were reported in two studies.[29,33] One stated that the patients included had ASA 3 or 4 and a body mass index of more than 40. However, no ASA number, sex, or age was reported in the article.[29] Mendes *et al.* did not report any background characteristics of the included patients.[33]

#### Description of the studies' risk of bias

Figure 2. shows the results of the risk of bias assessment. In all seven included studies, the cause and effect were clear. The majority of the studies measured outcomes in the same way and used appropriate statistical analyses. Several studies had limitations of follow-up and similarity in care and participants. None of the patients had multiple pre-and post-measurements.

#### Outcomes of the included studies

The outcomes of the included studies are described separately below.

#### Satisfaction

One study reported satisfaction as an outcome.[31] The summarised patient satisfaction with the anaesthetic consultation score out of 100 showed that patients in the PAC group

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> were more satisfied (mean difference, 2.10%; 95% confidence interval [CI], 0.51-3.70%; p=0.01).[31] There was no statistically significant difference between the two groups in the mean patient satisfaction with perioperative anaesthesia care score out of 5 after surgery (mean difference 0.01%, p=0.94).[31] The mean quality of recovery (QoR) score (range, 0-18) following anaesthesia on the first day of surgery was similar between the intervention (13.17±2.73) and control (13.31±2.65) groups (p=0.67).[31] The QoR measure is the patients' health-related quality of life.[35]

#### Anxiety

Three studies reported anxiety. [28,31,34] Two studies reported the visual analogue scale (VAS), one rated from zero (no anxiety) to ten (very high anxiety), [28] another used a 100 mm horizontal line with "not anxious at all" to "extremely anxious" [31] In one study, the median VAS anxiety score was 3(0-5) in the intervention group and 5(2-8) in the control group (p=0.0038).[28] In another study, there were no significant differences between the control and intervention groups for levels of anxiety (VAS), surgery (26 vs. 25, respectively, p=0.12), and anaesthesia (20 vs. 19, respectively, p=0.60).[31] The median Multiple Affect Adjective Check List (MAACL) score with possible range scores from 0 to 21 (higher scores indicating greater levels of anxiety) was 3 (0-9) in the intervention group and 6.5 (2-12) in the control group (p=0.0053).[28] The differences in the State-Trait Anxiety Index (STAI) score, which is composed of 40 questions rated on a 4-point Likert scale, was 1.51, 95% CI: 1.02–2.02%, p=0.0051).[34] The results on anxiety in these two studies were significant. However, Kamau et al. [34] found no differences when they examined anxiety and the influences of sex, duration of hospital stay, and prior anaesthesia experience. Mortality

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One study reported the mortality rates.[29] Patients attending the High Dependency Unit (HDU), Intensive Care Unit (ICU), and Post-anaesthesia Care Unit (PACU) following complex orthopaedic surgery had a significant reduction in mortality rate after being assessed in the PAC, from 18 (6.1%) of 298 patients before to 14 (1.2%) of 1147 patients after *p*=0.001.[29] *Cancellation rate* 

Three studies reported a reduced cancellation rate following the establishment of a PAC.[30,32,33] One of the included studies had 316 (2.0%) cancellations for medical reasons before the introduction of PAC, and 79 (0.9%) after, and a difference of 1.02% (95% CI, 0.31-1.31%). After adjustment, the odds ratio was 0.7 (95% CI, 0.5–0.9%).[30] The overall cancellation of surgery was reduced from 1027 (6.3%) to 393 (4.6%) following surgery, and a difference of 0.9% (95% CI, 0.3–1.0%) when patients were assessed in PAC.[30] Mendes et al.[33] found a decrease in overall cancellations from year 1 (39.3%) to year 4 (15.9%),  $p \le a$ 0.05. There were 469 (number of cancellations)/10639 (number of surgeries performed) due to medical reasons in the first year of this study. The following year, a considerable increase above the baseline values in the intervention group was observed, followed by a progressive decrease in the last year with 391 (number of cancellations)/10397 (number of surgeries performed).[33] Farasatkish et al. reported that of the 1716 patients studied, 15.1 % of cases cancelled in the two groups. The cancellation rates in the control group were 146/866 (16.8%), and the cancellation rate in the intervention group was 113/850 (13.29%) p=0.046. The most common reason for cancellation was incomplete medical work-up 51/146 (35%) in the control group and 32/113 (28%) in the intervention group, p=0.03).[32] Lee et al. found similar rates for surgery being cancelled on the scheduled date for the intervention group compared to the control group (2.3% vs. 3.4%, *p*=0.75).[31]

#### Costs and willingness to pay

Two studies reported the costs.[29,31] One study reported a total saving of £ 486.62 per patient after establishing a PAC.[29] Another study reported a significantly lower preoperative cost per patient in the intervention group compared to the control group (mean difference, \$ 463; 95% CI, -\$648 to -\$278 per patient, p<0.01).[31] However, the mean difference in the total perioperative treatment cost was not significant, even after adjusting for cancellation on the day of surgery costs.[31] The intervention group patients were willing to pay (WTP) significantly more than the median WTP (US \$13) for a clinic consultation at the PAC than the control group.[31]

#### Length of stay

The length of stay was reported in four studies.[29-31,33] Mendes *et al.*[33] found a significant decrease in mean hospital stay for patients from 6.2 to 5.0 days ( $p \le 0.001$ ) during the four years of this study. Van Klein *et al.*[30] found that the total admission time significantly decreased from a mean of 8.8 days (before) and a mean of 8.1 days (after) and 0.92 (0.90–0.94). After adjusting for age, sex, and introduction date of PAC this difference was 0.92 (0.90–0.94).[30] Kamal *et al.*[29] found a significant reduction in the length of stay in the high dependency unit from 2.1 days to 1.6 d (p=0.01), and in the intensive care unit from 2.3 days to 1.9 days (p=0.01). In the last study, no significant changes were found in the median duration of postoperative stay between the intervention and control groups.[31] *Organisation planning and efficiency* 

Organisation planning and efficiency have been reported in two studies.[29,33] One study found statistically significant changes in the reduction of unplanned admissions to the PACU (65/298 [22%], 111/1147 [10%], p=0.001), ICU (4/298 [1.3%], 4/1147 [0.4%], p=0.01), and HDU (4/298 [1.34%], 20/1147 [1.7%], p=0.01) after implementing a PAC.[29] The planned

admissions in the ICU (4/298 [1.3%], 18/1147 [1.6%], p=0.01), and HDU (14/298 [4.7%], 85/1147 [7.4%], p=0.1) increased after implementing a PAC.[29] The number of PAC evaluations increased from year 1, 4704 to year 4, 13990 (p≤ 0.001).[33] The number of outpatient procedures increased from 2170 (year 1) to 1943 (year 4) (p≤0.001), and the inpatient procedures decreased from 9556 (year 1) to 8449 (year 4), (p≤ 0.001).[33]

#### DISCUSSION

This systematic review summarises the effectiveness of PACs in improving quality and patient safety in general hospitals and determines the gaps in existing knowledge for future research. Seven studies that met the inclusion criteria were included. We present the main results and infer the implications for research and practice in the following text.

Cancellation on the day of surgery has undesirable effects on both the patients and the hospital system.[14] Thus, studies have found that late patient-related cancellations could totally or partially be prevented,[36] if they were addressed during preoperative evaluations.[14,15] This is confirmed by several studies in this systematic review that found a reduction in surgery cancellation after implementing a PAC.[30,32,33] However, Lee *et al.* found no significant changes between the intervention and control groups.[31] Mendes and colleagues found that the number of cancellations for medical reasons after PAC implementation decreased in the first year of implementation. In the second and third years, they were higher before the number dropped to below baseline.[33] These conflicting findings might show that hospitals operate in a specific context, with unique populations, processes, and microsystems, which may encounter unique obstacles making implementation difficult. Patient-focused interventions need to consider barriers,

facilitators, and interrelationships between systems, staff, and interventions to increase the likelihood of sustainable success.[37] In addition, Kamau *et al.* also indicated that PACs lead to more planned admissions to the ICU, HDU, and PACU, which is more predictable for patients, staff, and administrations.[34]

Another main finding of this systematic review was a significant reduction in the length of hospital stay following patients' examination in a PAC; however, a small number of studies with low quality were considered. Nevertheless, similar results were found in another systematic review claiming that perioperative systems support hospitals to address the expected growth in the number and complexity of surgical procedures being performed.[15] However, Lee *et al.* indicated that the reason for the reduced length of hospital stay was the mean duration of stay before surgery in the intervention group.[31] This indicates that when patients are examined in the PAC and well prepared with information, consultations, and tests, they do not need to be hospitalised until the day of surgery. A survey focusing on patients operated showed that if they had a choice, 75% do not wish to be admitted to the hospital until the same day of operation. One of the main reasons was to spend less time in the hospital.[38] However, an updated systematic review on the effectiveness of nurse-led preoperative assessment services for elective surgery found that the included articles had a reduced length of stay. The included studies had low methodological quality, and therefore, the authors could not conclude that this service leads to reduced length of hospital stay.[16]

The evidence from this systematic review is insufficient to conclude whether patients have reduced anxiety when assessed using PAC. The included studies used different instruments to measure the levels of anxiety, and the results could not be pooled. In addition, previous

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studies have shown that anxiety levels were higher in women.[39] Seventy-eight per cent of the participants were women in one of the included studies in this systematic review and might result in a bias in this study.[34] Anxiety was also statistically higher in patients who underwent general anaesthesia than in those who underwent regional anaesthesia.[40] The included studies on anxiety included both patients with general and regional anaesthesia, which might also be biased. Furthermore, the patients included in this review had both former surgical experience and no experience with surgery. However, studies have shown that former experience with anaesthesia and surgery reduces the risk of preoperative anxiety.[41]

Assessment of PAC was significantly associated with reduced mortality following complex orthopaedic surgery.[29] Previously published retrospective studies found similar results, but with other types of surgery.[42,43] A Danish study found that deaths attributed to anaesthesia were associated with insufficient or inadequate preoperative evaluation.[7] Furthermore, a previous study pointed out that the risk factors are not only patient-related but also organisation-related,[8] and that some hospitals have perioperative care and teams that are better at identifying and rescuing perioperative complications.[44,45] However, Blitz *et al.* argued that PAC should focus on early patient engagement strategies, interdisciplinary team communication, detailed perioperative care plans, and patient documentation in the electronic health record. This record should be open for review by the perioperative team to preserve patient information and safety. The value of a PAC lies in its ability to improve the quality of the perioperative process by designing a more robust system for preoperative assessment and preparation.[42] The importance of safety in

anaesthesia is a vital component in anaesthesia practice, and the use of PACs contributes to this critical area.

#### Strengths and limitations of the study

Most review steps were performed in duplicate or independently by two researchers, and agreement was reached in a consensus meeting. However, grey literature, such as government and institutional documents, was not included and might be a limitation to this study. Since countries have different organisational structures in their healthcare systems, we did not set inclusion criteria concerning who performed the patient's preoperative assessment. However, the European Society of Anaesthesiology guidelines recommend that anaesthesiologists complete the preoperative assessment, while trained nurses or anaesthesia trainees perform the screening.[8] A preoperative evaluation performed by an internist has been associated with increased length of stay and increased postoperative mortality.[46] This systematic review's results were possibly affected by the heterogeneity in the types of staff performing the preoperative assessment.

We opted to include only the studies with the highest internal validity. Thus, we excluded several retrospective studies. Nonetheless, the remaining studies' risk of bias was fairly high, and they were heterogeneous. As a result, meta-analyses were not statistically appropriate.[25] The included studies' designs could not rule out selection bias and confounding, and the strength of the evidence should be assessed cautiously. Many studies did not make adjustments for several confounders, which could be responsible for the observed effects. Several studies lacked descriptions of the methods used and the patients included, which lowered the transparency. It is not very reassuring that many such studies were unable to deliver more thorough evidence to guide practice and should be assessed

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cautiously. The results are relevant to health care services, which should focus on the wellbeing and safety of the patients.

#### Implications for future research and practice

This systematic review identified the ambiguity in the PAC interventions offered to the intervention group. In many studies, it was evident that the methods used in these studies were not always clearly described, and high-quality research is needed in this field. The included studies in this review did not contain any results of reduced preoperative tests, such as blood tests, on patients before surgery when patients attended the PAC, [47,48] and earlier surgical room entry time for patients assessed in PACs, [49,50] similar to previous retrospective studies. Other implications for future research might be the organisation structure of different PACS and their functioning. The use of technology, such as streaming services, facilitates different types of patient groups and might be more important with the appearance of Covid-19 in reducing human contact and spread of the virus.

#### CONCLUSION

This systematic review suggests that PAC use reduces the length of hospital stay, and the majority of the studies had reduced the cancellation rate in hospitals. These findings are an essential contribution to the current evidence in this field. In addition to further research in this field, the demand for increased high-quality studies to capture robust data describing the quality of care and clinical outcomes for patients requiring anaesthesia. This step demands increased focus and funding for this specific area of health services research and could, therefore, lead to new implementations of PAC's in health care services and further develop patient safety in perioperative care.

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The authors declare that they have no conflict of interest.

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#### Author statement

EWK, MF, AO and TOT: Study design

EWK, MF: Search and screening of the articles

EWK: Data extraction

MF, AO, RB and TOT: Control of data extraction

EWK, MF, AO and RB: Quality assessment of the included articles

EWK: Drafting of the manuscript

MF, AO, RB, and TOT: Contribution to and review of the final version of the manuscript

MF, AO, RB and TOT: Supervised the study

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From: Moher D, Liberati A, Tetzlaff J, Altman DG, The PRISMA Group (2009). Preferred Reporting Items for Systematic Reviews and Meta-Analyses: The PRISMA Statement. PLoS Med 6(7): e1000097. doi:10.1371/journal.pmed1000097

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#### Figure 2: The risk of bias assessment



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1 ((pre-operativ\* or preoperativ\* or pre operativ\*) and (assessment\* or measurement\* or evaluat\*)).ti. (6182)

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# Reporting checklist for systematic review and meta-analysis.

Based on the PRISMA guidelines.

# Instructions to authors

Complete this checklist by entering the page numbers from your manuscript where readers will find each of the items listed below.

Your article may not currently address all the items on the checklist. Please modify your text to include the missing information. If you are certain that an item does not apply, please write "n/a" and provide a short explanation.

Upload your completed checklist as an extra file when you submit to a journal.

In your methods section, say that you used the PRISMAreporting guidelines, and cite them as:

Moher D, Liberati A, Tetzlaff J, Altman DG, The PRISMA Group. Preferred Reporting Items for Systematic Reviews and Meta-Analyses: The PRISMA Statement

			Page	
		Reporting Item	Number	
Title		4		
	<u>#1</u>	Identify the report as a systematic review, meta-analysis, or both.	1	
Abstract				
Structured summary	<u>#2</u>	Provide a structured summary including, as applicable: background; objectives; data sources; study eligibility criteria, participants, and interventions; study appraisal and synthesis methods; results; limitations; conclusions and implications of key findings; systematic review registration number	2	
Introduction				
Rationale	<u>#3</u>	Describe the rationale for the review in the context of what is already known.	3-5	
	For p	peer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml		
1 2 3 4 5	Objectives	<u>#4</u>	Provide an explicit statement of questions being addressed with reference to participants, interventions, comparisons, outcomes, and study design (PICOS).	5
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6 7	Methods			
8 9 10 11 12	Protocol and registration	<u>#5</u>	Indicate if a review protocol exists, if and where it can be accessed (e.g., Web address) and, if available, provide registration information including the registration number.	5
13 14 15 16 17 18 19	Eligibility criteria	<u>#6</u>	Specify study characteristics (e.g., PICOS, length of follow- up) and report characteristics (e.g., years considered, language, publication status) used as criteria for eligibility, giving rational	6
20 21 22 23 24 25 26	Information sources	<u>#7</u>	Describe all information sources in the search (e.g., databases with dates of coverage, contact with study authors to identify additional studies) and date last searched.	6
27 28 29 30 31	Search	<u>#8</u>	Present full electronic search strategy for at least one database, including any limits used, such that it could be repeated.	5 + APPENDIX
32 33 34 35 36 37	Study selection	<u>#9</u>	State the process for selecting studies (i.e., for screening, for determining eligibility, for inclusion in the systematic review, and, if applicable, for inclusion in the meta-analysis).	6
38 39 40 41 42 43	Data collection process	<u>#10</u>	Describe the method of data extraction from reports (e.g., piloted forms, independently by two reviewers) and any processes for obtaining and confirming data from investigators.	6
44 45 46 47 48 49	Data items	<u>#11</u>	List and define all variables for which data were sought (e.g., PICOS, funding sources), and any assumptions and simplifications made.	5-6
50 51 52 53 54 55 56	Risk of bias in individual studies	<u>#12</u>	Describe methods used for assessing risk of bias in individual studies (including specification of whether this was done at the study or outcome level, or both), and how this information is to be used in any data synthesis.	6
57 58 59 60	Summary measures	<u>#13</u> For pe	State the principal summary measures (e.g., risk ratio, difference in means). eer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml	N/A

1 2 3 4 5	Planned methods of analyis	<u>#14</u>	Describe the methods of handling data and combining results of studies, if done, including measures of consistency (e.g., I2) for each meta-analysis.	7
6 7 8 9 10	Risk of bias across studies	<u>#15</u>	Specify any assessment of risk of bias that may affect the cumulative evidence (e.g., publication bias, selective reporting within studies).	11-12
11 12 13 14 15 16	Additional analyses	<u>#16</u>	Describe methods of additional analyses (e.g., sensitivity or subgroup analyses, meta-regression), if done, indicating which were pre-specified.	N/A
17 18	Results			
19 20 21 22 23	Study selection	<u>#17</u>	Give numbers of studies screened, assessed for eligibility, and included in the review, with reasons for exclusions at each stage, ideally with a <u>flow diagram</u> .	8
24 25 26 27 28	Study characteristics	<u>#18</u>	For each study, present characteristics for which data were extracted (e.g., study size, PICOS, follow-up period) and provide the citation.	N/A
29 30 31 32	Risk of bias within studies	<u>#19</u>	Present data on risk of bias of each study and, if available, any outcome-level assessment (see Item 12).	11-12
33 34 35 36 37 38 39	Results of individual studies	<u>#20</u>	For all outcomes considered (benefits and harms), present, for each study: (a) simple summary data for each intervention group and (b) effect estimates and confidence intervals, ideally with a forest plot.	14-17
40 41 42 43 44	Synthesis of results	<u>#21</u>	Present the main results of the review. If meta-analyses are done, include for each, confidence intervals and measures of consistency.	14-17
45 46 47 48	Risk of bias across studies	<u>#22</u>	Present results of any assessment of risk of bias across studies (see Item 15).	N/A
49 50 51 52	Additional analysis	<u>#23</u>	Give results of additional analyses, if done (e.g., sensitivity or subgroup analyses, meta-regression [see Item 16]).	N/A
53 54	Discussion			
55 56 57 58	Summary of Evidence	<u>#24</u>	Summarize the main findings, including the strength of evidence for each main outcome; consider their relevance	18-20
59 60		For p	eer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml	

Page	37 of 36		BMJ Open	
1 2 2			to key groups (e.g., health care providers, users, and policy makers	
5 4 5 6 7 8	Limitations	<u>#25</u>	Discuss limitations at study and outcome level (e.g., risk of bias), and at review level (e.g., incomplete retrieval of identified research, reporting bias).	20-21
9 10 11 12	Conclusions	<u>#26</u>	Provide a general interpretation of the results in the context of other evidence, and implications for future research.	22
13 14	Funding			
15 16 17 18 19	Funding	<u>#27</u>	Describe sources of funding or other support (e.g., supply of data) for the systematic review; role of funders for the systematic review.	23
20 21 22	None The PRISM	A check	klist is distributed under the terms of the Creative Commons At	tribution
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# Effectiveness of pre-anaesthetic assessment clinic: a systematic review of randomised and non-randomised prospective controlled studies

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<b>Primary Subject Heading</b> :	Anaesthesia
Secondary Subject Heading:	Surgery, Nursing, Health services research
Keywords:	Adult anaesthesia < ANAESTHETICS, Health & safety < HEALTH SERVICES ADMINISTRATION & MANAGEMENT, Quality in health care < HEALTH SERVICES ADMINISTRATION & MANAGEMENT, Adult surgery < SURGERY
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1	Effectiveness of pre-anaesthetic assessment clinic: a systematic review of randomised and
2	non-randomised prospective controlled studies
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4	Eirunn Wallevik Kristoffersen <sup>1,2</sup> *, Anne Opsal <sup>1</sup> , Tor Oddbjørn Tveit <sup>1,2,3</sup> , Rigmor C Berg <sup>4,5</sup> ,
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2 3	24	ABSTRACT
4 5	25	Objectives: The aim of this systematic review was to examine the effectiveness of pre-
6 7	26	anaesthetic assessment clinics (PACs) implemented to improve quality and patient safety in
8 9	27	perioperative care.
10 11	28	Design: Systematic review.
12 13	29	Data sources: The electronic databases CINAHL Plus with Full Text (EBSCOhost), Medline,
14 15	30	and Embase (OvidSP) were systematically searched from 1st April, 1996 to 4th February,
15 16 17	31	2021.
17	32	Eligibility criteria: The main inclusion criterion was that the study, using empirical
19 20	33	quantitative methods, addressed the effectiveness of PACs.
21 22	34	Data extraction and synthesis: Titles, abstracts, and full texts were screened by a team of
23 24	35	three authors. Risk of bias was assessed using the Joanna Briggs Institute critical appraisal
25 26	36	checklist for quasi-experimental studies. Data extraction was performed by one author and
27	37	checked by four other authors. Results were synthesised narratively owing to the
28 29	38	heterogeneity of the included studies.
30 31	39	<b>Results:</b> Seven prospective controlled studies, on the effectiveness of PACs, were included.
32 33	40	Three studies reported a significant reduction in the length of the hospital stay, and two
34 35	41	studies reported a significant reduction in cancellation of surgery for medical reasons when
36 37	42	patients were seen in the PAC. In addition, the included studies presented mixed results
38	43	regarding anxiety in patients. Most studies had a high risk of bias.
40	44	Conclusion: This systematic review demonstrated a reduction in the length of hospital stay
41 42	45	and cancellation of surgery when the patients had been assessed in the PAC. There is a need
43 44	46	for high-quality prospective studies to gain a deeper understanding of the effectiveness of
45 46	47	PACs.
47 48	48	PROSPERO registration number: CRD42019137724
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50 51	50	Keywords: pre-anaesthetic assessment clinic, preoperative care, quality, safety, systematic
52 53	51	review
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- 3 4	58	Strengths and Limitations of this study
5	59	An extensive database search was conducted with no limitations on outcomes and
6 7	60	the type of pre-anaesthetic assessment clinic.
8 9	61	Only randomised or non-randomised prospective controlled studies were included.
10 11	62	• The Joanna Briggs Institute Critical appraisal checklist for quasi-experimental studies
12 13	63	was used.
14 15	64	• The included studies were heterogeneous and had high risk of bias, which is a major
16	65	limitation of this review.
17 18	66	
19 20	67	
21 22	60	A normalization is an evid in success the second structure where is a single sized shows a short
23 24	68	Anaestnesia is crucial in surgery. However, it may activate physiological changes that
24 25	69	increase morbidity and mortality, [1] depending on the patients' preoperative health status
26 27	70	and age. [2] Hospitals treat patients with complex, comorbid healthcare problems
28 29		
30 31	71	undergoing progressively extensive surgeries and interventions. [3,4] To ensure the quality
32	72	and safety of anaesthesia and surgery, precise knowledge of the clinical characteristics of
34	73	patients undergoing surgery is critical to the perioperative treatment plan. [2] Over the past
35 36		
37 38	74	50 years, perioperative mortality, including anaesthesia-related mortality, has declined,
39 40	75	which is significant in developed countries, [1,5] mainly due to new anaesthetics, improved
41 42	76	monitoring equipment and training, availability of recovery rooms, and improved airway
43		
44 45	77	management. [4] However, an Australian study reported that 14% of anaesthetic-surgical
46 47	78	complications and 39% of deaths attributed to anaesthesia were associated with insufficient
48 49	79	and/or inadequate preoperative evaluation [6] A Danish retrospective investigation showed
50 51	, ,	
52	80	that the deaths among patients undergoing surgery could be prevented by thorough
55 54	81	preoperative evaluation, [7] indicating that risk factors are both patient-and surgery-related
55 56	82	and linked to organisational structures [8] Future efforts should improve preoperative
57 58	52	
59 60		

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anaesthesia safety, [9] by improving planning and preparation for elective procedures and interventions.

Today, an increasing number of pre-anaesthesia assessment clinics (PACs) support hospitals internationally in handling the rising number of patients and complexity of surgical procedures. [10] The design of PACs differs crtically based on location, organisational structure, timing, and patient groups. They primarily function as a service unit for surgeons, patients, and the anaesthetic team. [11] The PAC consultation, by the anaesthesiologist, anaesthesia nurse, or both, is a globally recognised evaluation method and optimises the patients' medical condition prior to surgery and anaesthesia. [2] Thus, it is essential for secure anaesthetic practice since it detects anaesthesia-related risk factors and high-risk patients, improves patient outcomes, prepares the patient physically and psychologically for anaesthesia, and ensures the patient's most favourable condition for surgery and anaesthesia. [12-14] This is primarily performed by interviewing and examining the patient, reviewing previous medical, surgical and anaesthesia issues, detailed description of current medication, and provisions for obtaining and reviewing preoperative tests. [11] PACs also lead to increased communication between healthcare providers and coordination with postoperative care. [15,16] Due to well-prepared patients and staff, several researchers posit that with PAC, the number of surgical cancellations, length of hospital stay, and mortality rate have reduced, and tests are minimised. [8,17,18] Others assert that patients feel less anxious regarding the subsequent anaesthetic and surgical processes and are highly satisfied with the service with PAC consultations. [17,19,20]

As Turunen et al. stated, research on PACs is scarce regarding costs, financial savings, the impact on patient safety and quality of care, accuracy of operative patients, and effect on

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.06 preoperative nursing levels. [21] Survey results indicate that anaesthesiologists perceive day .07 of surgery delays due to missing information as common, even with PAC consultations. [22] This systematic review aimed to examine the effectiveness of PACs in improving quality and .08 .09 patient safety in preoperative care. Further, we aimed to determine the gaps in existing .10 knowledge for future research. **METHODS** .11 .12 Our systematic review followed the guidelines in the Cochrane Handbook for Systematic .13 Reviews of Interventions [23] and was reported in accordance with the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines. [24] The protocol was .14 .15 registered in PROSPERO: CRD42019137724. [25] .16 The two review questions were: 1. Is PAC effective in improving patient satisfaction and safety, while reducing anxiety? .17 .18 2. Is PAC effective in reducing cancellation rate, cost, and improving efficiency? .19 Search strategies .20 We performed a scoping search in different databases to identify the key terms. [26,27] The .21 final search was planned and conducted in close collaboration with a university librarian. On .22 11th September, 2018 we searched CINAHL Plus with Full Text (EBSCOhost), Medline, and .23 Embase (OvidSP), which was updated on 3th February 2020 and 4th February, 2021. .24 Considering the lack of subject headings (e.g., MeSH) for PAC, we combined text words such .25 as preanaesthesia, nurse, surgery, anaesthesia, preoperative, assessment, measurement, .26 evaluate, preadmission, centre, clinic, ward, unit, and outpatient. The searches are detailed .27 in Appendix 1. The search mode in CINAHL was Boolean/Phase, which supports Boolean .28 searching or exact phrase searching. For comprehensiveness, we used both the truncation

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2 3	129	and proximity operators. We limited the search to 1996, the year one of the first known
4 5 6	130	articles in this area was published. [28] Complementary methods to identify studies included
7 8	131	following up on citations via Scopus, scanning the reference lists of relevant papers and
9 10 11	132	included articles, and checking for relevant studies in clinical trials. [26]
12 13	133	Eligibility criteria
14 15 16	134	The main inclusion criterion was that the study, using empirical quantitative methods.
16 17 18	135	addressed the effectiveness of PACs. Specific eligibility criteria were: (a) published in English
19 20	136	or Scandinavian language (h) scientific nublication of original research (c) reported the
21 22 23	137	outcomes of PAC (d) PAC consultation with the patient present (e) randomised or non-
24 25	120	randomised prospective controlled studies, and (f) newly established BAC. We excluded: (a)
26 27 28	120	aditorials discussions and conference abstracts (b) reviews (c) instrument testing (d)
28 29 30	135	studios on childron, and (o) retrospective studios
31 32	140	Study selection
33 34	141	Study selection
35 36	142	All references identified in the search were transferred to EndNoteX9, where the duplicates
37 38 39	143	were removed. Subsequently, all unique references were transferred to the Covidence
40 41	144	screening tool. [29] Study eligibility was ascertained independently, first at the title and
42 43	145	abstract level, and subsequently at the full text. Three of the authors screened all the
44 45 46	146	articles (EWK, AO, MF). Inclusion was determined by consensus, and disagreements were
47 48	147	resolved by consulting RCB and TOT.
49 50 51	148	
52 53	149	Quality assessment
54 55 56	150	Risk of bias in studies were assessed using design specific checklists. Given the
57 58	151	methodological similarity of the studies, only the Joanna Briggs Institute Critical appraisal
59 60	152	checklist for quasi-experimental studies was used. [30] Author EWK performed the risk of

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2 3 4	153	bias assessment, and RCB confirmed its accuracy. Disagreements were resolved through
5 6 7	154	discussion with MF and AO. Each of the nine checklist questions was answered no, yes,
7 8 9	155	unclear (or not applicable).
10 11	156	Data extraction and analysis
12 13 14	157	Author EWK extracted data from each study onto a pre-designed Excel spreadsheet. All the
15 16 17	158	authors confirmed the accuracy, consistency, and completeness of the extracted data that
17 18 19	159	included publication details, study design, setting, and characteristics of the patients,
20 21	160	interventions, comparisons, and outcome (PICO). We requested information on the missing
22 23 24	161	data; however, received no response from the authors. If the PICO elements were
25 26	162	sufficiently similar and statistical data available, we intended to conduct meta-analyses.
27 28 29	163	However, the extracted data revealed substantial heterogeneity. Therefore, we performed
30 31	164	narrative synthesis, describing and comparing the main findings from the included studies,
32 33 34	165	and discussing their methodological strengths and weaknesses.
35 36	166	Patient and Public Involvement
37 38 30	167	Patients and/or the public were not involved in the design, conduct, reporting, or
40 41	168	dissemination plans of this research. However, the project is initiated by health
42 43	169	professionals.
44 45 46	170	
47 48	171	RESULTS
49 50 51	172	Figure 1. provides details of the study selection process. A total of 2,981 records were
52 53	173	identified in the final search (2021). After removing duplicates, we screened 2,058 records
54 55 56	174	based on the title and abstract; 179 records passed the full-text screening. We included
57 58	175	seven studies that met the inclusion criteria.
59 60	176	Overall characteristics of the studies

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The seven studies are listed in Table 1. They were all in English and published between 2000–2017, with data collected in the years 1997–2015 (one study did not report data collection information). [31] Based on inclusion criteria, all were prospective controlled studies; however, no RCTs were found. There was one controlled before-after study. [32] The rest six studies had control groups; assessments followed PAC implementation, without baselinee assessments. There were three 2-group non-parallel after-only studies, [33-35] and three 2-group parallel after-only studies [31]; one had a matched control group [36] and one had three follow-up assessments of one arm. [37] One study had only cancellation rate as prospective data. [33] The studies included 77,411 patients. 

### **Table 1: Description of included studies**

Author, Year, Country	Study design	Sampling time	Population	Intervention	Comparison	Outcomes
Farasatkish,2009[34] Iran	2-group after study	May 2007 through August 2007	N=1716, open- heart surgery, ASA class III-IV	Pre-anaesthesia consultation clinic (3- 10 days before surgery)	Usual care (within 24 h of surgery)	Cancellations
Kamal,2011[35] England	2-group after study	April 2005 through April 2009	N=1445, complex elective orthopaedic surgery, ASA class III-IV	Preoperative anaesthetic assessment clinic (timing not stated)	Usual care (day of surgery)	Admissions, length of stay, mortality, cost
Kamau,2017[32] Kenya	CBA	August 2000, April 2001, November 2001	N=51, elective non-cardiac surgery, ASA class not stated	Pre-anaesthesia clinic consultation (≥48 h before surgery)	Usual care (day before surgery)	Anxiety (STAI score)
Klopfenstein,2000[31] Switzerland	2-group after study (parallel)	No data	N=40, elective endoscopic urological surgery, ASA class I-III	Pre-anaesthetic consultation (1-2 weeks before surgery)	Usual care (the evening before surgery)	Anxiety (MAACL, VAS)
Lee,2012[36] China	2-group after study (parallel)	March 2007 through November 2009	N=352, elective surgery, ASA class I-IV	Anaesthesia consultation clinic (≤3 months before surgery)	Usual care (the evening before surgery)	Quality of recovery score, cost, cancellations, length of stay, satisfaction, anxiety (VAS),

Mendes,2005[37] Brazil	2-group after study (parallel)	April 2007 through June 2007	N=52254, surgery, ASA class not stated	Preoperative outpatient evaluation clinic (timing not	Usual care (timing not stated)	willingness to pay (WTP) Cancellations, length of stay
van Klei,2002[33] The Netherlands	2-group after study	November 2012	N=21553, elective surgery, ASA class not stated	Preoperative outpatient evaluation clinic (average 3 weeks before surgery)	Usual care (day before surgery)	Cancellations

ASA: American Society of Anesthesiology; CBA: controlled before-after; MAACL: Multiple Affect Adjective Check List; STAI: State-Trait Anxiety Inventory; VAS: visual analogue scale; WTP: willingness to pay

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190	Considering the intervention, PACs in all studies comprised an outpatient service whereby
191	patients were examined for medical conditions important for anaesthesia and informed
192	regarding expectations on the day of surgery. Nevertheless, the terminology used for PACs
193	varied, they served different surgical specialities, and conducted pre-anaesthesia
194	consultation from $\geq$ 48 h to $\leq$ 3 months before the surgery. The settings included university
195	hospital (n=3), [32,36,37] teaching hospital (n=1), [33] medical centre(n=1), [34] and general
196	hospital (n=1) [35]; one study did not specify the context.[31] The staff conducting the pre-
197	anaesthesia consultation also varied: in five studies, it was the anaesthesiologists, [31,33,35-
198	37] in the other studies it was (also) the orthopaedic senior house officer, [35] the
199	consultant or resident, [32] or the physician. [34] In three studies, nurses were part of the
200	team. [33,35,36] The comparison in all studies was usual care, which generally involved a
201	preoperative anaesthetic evaluation of the admitted patients the day before the surgery.
202	
203	Of the 77,411 patients in the studies, 9,626 and 15,531 patients were in the intervention and
204	control groups, respectively. One study did not specify the number of patients in the
205	intervention and control groups, but only the total number of surgeries performed. [37] Five
205 206	intervention and control groups, but only the total number of surgeries performed. [37] Five studies reported data for sex, showing that 51% of the patients were women and 49% were
205 206 207	intervention and control groups, but only the total number of surgeries performed. [37] Five studies reported data for sex, showing that 51% of the patients were women and 49% were men (12,129 vs. 11,583). [31-34,36] There were more women than men in both the
205 206 207 208	intervention and control groups, but only the total number of surgeries performed. [37] Five studies reported data for sex, showing that 51% of the patients were women and 49% were men (12,129 vs. 11,583). [31-34,36] There were more women than men in both the intervention (4,345 vs. 4,134) and the control groups (7,784 vs. 7,449). Five studies reported
205 206 207 208 209	intervention and control groups, but only the total number of surgeries performed. [37] Five studies reported data for sex, showing that 51% of the patients were women and 49% were men (12,129 vs. 11,583). [31-34,36] There were more women than men in both the intervention (4,345 vs. 4,134) and the control groups (7,784 vs. 7,449). Five studies reported data for age showing that all the patients were over 20 years old [31-34,36] and four studies
205 206 207 208 209 210	intervention and control groups, but only the total number of surgeries performed. [37] Five studies reported data for sex, showing that 51% of the patients were women and 49% were men (12,129 vs. 11,583). [31-34,36] There were more women than men in both the intervention (4,345 vs. 4,134) and the control groups (7,784 vs. 7,449). Five studies reported data for age showing that all the patients were over 20 years old [31-34,36] and four studies had grouped within the American Society of Anesthesiology (ASA) category. [31,34-36]
205 206 207 208 209 210 211	intervention and control groups, but only the total number of surgeries performed. [37] Five studies reported data for sex, showing that 51% of the patients were women and 49% were men (12,129 vs. 11,583). [31-34,36] There were more women than men in both the intervention (4,345 vs. 4,134) and the control groups (7,784 vs. 7,449). Five studies reported data for age showing that all the patients were over 20 years old [31-34,36] and four studies had grouped within the American Society of Anesthesiology (ASA) category. [31,34-36]
205 206 207 208 209 210 211 211	intervention and control groups, but only the total number of surgeries performed. [37] Five studies reported data for sex, showing that 51% of the patients were women and 49% were men (12,129 vs. 11,583). [31-34,36] There were more women than men in both the intervention (4,345 vs. 4,134) and the control groups (7,784 vs. 7,449). Five studies reported data for age showing that all the patients were over 20 years old [31-34,36] and four studies had grouped within the American Society of Anesthesiology (ASA) category. [31,34-36] The patients were scheduled to undergo a variety of surgeries, including orthopaedic,
205 206 207 208 209 210 211 212 212	intervention and control groups, but only the total number of surgeries performed. [37] Five studies reported data for sex, showing that 51% of the patients were women and 49% were men (12,129 vs. 11,583). [31-34,36] There were more women than men in both the intervention (4,345 vs. 4,134) and the control groups (7,784 vs. 7,449). Five studies reported data for age showing that all the patients were over 20 years old [31-34,36] and four studies had grouped within the American Society of Anesthesiology (ASA) category. [31,34-36] The patients were scheduled to undergo a variety of surgeries, including orthopaedic, [32,33,35,36] urology, [31-33,36] general, [32,33,36] heart, [34] gynaecology/obstetrics,

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2 3 4	214	[32,33,36] vascular surgery, [33] ophthalmology, [33] maxillofacial/dental surgery,
5 6 7 8 9	215	[32,33]neurological surgery, [33] while one did not specify the type. [37] In five studies, the
	216	type of anaesthesia was not specified, [32-35,37] and two studies reported patients for
10 11	217	general and/or regional supplement. [31,36]
12 13 14	218	
15 16	219	The patients included had previous anaesthetic experience in one study, [31] previous and
17 18 19	220	no previous anaesthetic experience in another, [32] and five studies did not report this data.
20 21	221	[33-37] Limited background characteristics of the patients were reported in two studies.
22 23 24	222	[35,37] One stated that the patients included had ASA 3 or 4 and a body mass index >40;
24 25 26	223	however, no ASA number, sex, or age was reported. [35] Mendes et al. did not report any
27 28 29	224	background characteristics of the included patients. [37]
30 31	225	Description of risk of bias in the studies
32 33 24	226	Figure 2. shows the results of the risk of bias assessment. In all seven included studies, the
34 35 36	227	cause and effect were clear. The majority of the studies measured outcomes similarly and
37 38	228	used appropriate statistical analyses. Several studies had limitations of follow-up and
39 40 41	229	similarity in care and participants. None of the patients had multiple pre-and post-
42 43	230	measurements.
44 45	231	Outcomes of the included studies
46 47 48	232	The outcomes of the included studies are each described separately below.
49 50	233	Satisfaction
52 53	234	One study reported satisfaction as an outcome. [36] The summarised patient satisfaction
54 55	235	with the anaesthetic consultation score out of 100 showed that patients in the PAC group
50 57 58	236	were more satisfied (mean difference, 2.10%; 95% confidence interval [CI], 0.51–3.70%;
59 60	237	p=0.01). [36] There was no statistically significant difference between the two groups in

mean patient satisfaction with perioperative anaesthesia care score out of 5 after surgery (mean difference 0.01%, p=0.94). [36] The quality of recovery (QoR) measure is the patients' quality of recovery score. [38] The mean QoR score (range, 0–18) following anaesthesia on the first day after surgery was similar between the intervention (13.17±2.73) and control (13.31±2.65) groups (*p*=0.67). [36]

Anxiety

> Three studies reported anxiety. [31,32,36] Two studies reported the visual analogue scale (VAS), one rated from zero (no anxiety) to ten (very high anxiety), [31] another used a 100 mm horizontal line with "not anxious at all" to "extremely anxious [36]" In one study, the median VAS anxiety score was 3 (0–5) in the intervention group and 5 (2–8) in the control group (p=0.0038). [31] In another study, there were no significant differences between the control and intervention groups for levels of anxiety (VAS), surgery (26 vs. 25, respectively, p=0.12), and anaesthesia (20 vs. 19, respectively, p=0.60). [36] The median Multiple Affect Adjective Check List (MAACL) score, with possible range of scores 0–21 (higher scores indicating greater levels of anxiety), was 3 (0–9) in the intervention group and 6.5 (2–12) in the control group (p=0.0053). [31] The differences in the State-Trait Anxiety Index (STAI) score, which comprising 40 questions rated on a 4-point Likert scale, was 1.51, 95% CI: 1.02-2.02%, p=0.0051). [32] The results on anxiety in these two studies were significant. However, Kamau et al. found no differences on examining anxiety and the influences of sex, duration of hospital stay, and prior anaesthesia experience. [32] Mortality One study reported the mortality rates. [35] Patients attending the High Dependency Unit

(HDU), Intensive Care Unit (ICU), and Post-anaesthesia Care Unit (PACU) following complex

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orthopaedic surgery had a significant reduction in mortality rate after being assessed in the PAC, from 18 (6.1%) of 298 patients before to 14 (1.2%) of 1147 patients after p=0.001. [35] Cancellation rate Four studies reported reduced cancellation rates following the establishment of a PAC. 

[33,34,36,37] One of the included studies had 316 (2.0%) cancellations for medical reasons before the introduction of PAC, and 79 (0.9%) after, and a difference of 1.02% (95% CI, 0.31-1.31%). After adjustment, the odds ratio was 0.7 (95% CI, 0.5–0.9%). [33] The overall cancellation of surgery reduced from 1027 (6.3%) to 393 (4.6%) following surgery, and a difference of 0.9% (95% CI, 0.3–1.0%) when patients were assessed in PAC. [33] Mendes et al. [37] found a decrease in overall cancellations from year 1 (39.3%) to year 4 (15.9%),  $p \le$ 0.05. There were 469 (number of cancellations)/10,639 (number of surgeries performed) due to medical reasons in the first year of this study. The following year, a considerable increase above the baseline in the intervention group was observed, followed by a progressive decrease in the last year with 391 (number of cancellations)/10397 (number of surgeries performed). [37] Farasatkish *et al.* reported that of the 1,716 patients studied, 15.1 % cancelled in the two groups. The cancellation rates in the control and intervention groups were 146/866 (16.8%) and 113/850 (13.29%) (p=0.046), respectively. The most common reason for cancellation was incomplete medical work-up; 51/146 (35%) in the control group and 32/113 (28%) in the intervention group (p=0.03). [34] Lee et al. found similar rates for surgery being cancelled on the scheduled date for the intervention group compared to the control group (2.3% vs. 3.4%, *p*=0.75). [36] 

Costs and willingness to pay

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284 Two studies reported the costs. [35,36] One study reported a total saving of £ 486.62 per 285 patient after establishing a PAC. [35] Another reported a significantly lower preoperative 286 cost per patient in the intervention group compared to the control group (mean difference, \$ 463; 95% Cl, -\$648 to -\$278 per patient, *p*<0.01). [36] However, the mean difference in the 287 288 total perioperative treatment cost was not significant, even after adjusting for cancellation 289 on the day of surgery costs. [36] The intervention group patients were willing to pay (WTP) 290 significantly more than the median WTP (US \$13) for a clinic consultation at the PAC than the control group. [36] 291

292 Length of stay

The length of stay was reported in three studies. [35-37] Mendes *et al.* [37] found a significant decrease in mean hospital stay of patients from 6.2 to 5.0 days ( $p \le 0.001$ ) during the four years of this study. Kamal *et al.* [35] found a significant reduction in the length of stay in the high dependency unit from 2.1 days to 1.6 d (p=0.01), and in the intensive care unit from 2.3 days to 1.9 days (p=0.01). In the last study, no significant changes were found in the median duration of postoperative stay between the intervention and control groups. [36]

300 Organisation planning and efficiency

Organisation planning and efficiency have been reported in two studies. [35,37] One study found statistically significant changes in the reduction of unplanned admissions to the PACU (65/298 [22%], 111/1147 [10%], p=0.001), ICU (4/298 [1.3%], 4/1147 [0.4%], p=0.01), and HDU (4/298 [1.34%], 20/1147 [1.7%], p=0.01) after implementing a PAC. [35] The planned admissions in the ICU (4/298 [1.3%], 18/1147 [1.6%], p=0.01), and HDU (14/298 [4.7%], 85/1147 [7.4%], p=0.1) increased after implementing a PAC. [35] The number of PAC evaluations increased from year 14,704 to year 413,990 (p≤ 0.001). [37] The number of

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3 4	308	outpatient procedures increased from 2,170 (year 1) to 1,943 (year 4) ( $p\leq 0.001$ ), and the
5 6 7	309	inpatient procedures decreased from 9,556 (year 1) to 8,449 (year 4), ( $p \le 0.001$ ). [37]
8	310	
9 10	311	DISCUSSION
11 12 13	312	This systematic review summarises the effectiveness of PACs in improving quality and
14 15 16 17 18	313	patient safety in general hospitals and determines the gaps in existing knowledge for future
	314	research. Seven studies met the inclusion criteria. Herein, we present the main results and
19 20	315	infer the implications for research and practice.
21 22 23	316	
24 25	317	Cancellation on the day of surgery has undesirable effects on both the patients and the
26 27 28	318	hospital system. [14] Studies have found that late patient-related cancellations could totally
29 30 31 32 33 34 35 36 37 38 39 40	319	or partially be prevented, [39] if addressed during preoperative evaluations. [14,17] This is
	320	confirmed by only three studies in this systematic review that found a reduction in surgery
	321	cancellation after implementing a PAC. [33,34,37] However, Lee <i>et al</i> . found no significant
	322	changes between the intervention and control groups. [36] Mendes <i>et al.</i> found that the
	323	number of cancellations for medical reasons after PAC implementation decreased in the first
41 42	324	year of implementation. In the second and third years, they were high before the number
43 44 45	325	dropped to below baseline. [37] These conflicting findings might show that hospitals operate
46 47	326	in specific contexts, with unique populations, processes, and microsystems, encountering
48 49 50	327	unique obstacles, making implementation difficult. Patient-focused interventions should
50 51 52	328	consider barriers, facilitators, and interrelationships between systems, staff, and
53 54	329	interventions to increase the likelihood of sustainable success. [40] Addition, Kamau et al.
55 56 57	330	indicated that PACs lead to more planned admissions to the ICU, HDU, and PACU, which is
58 59 60	331	more predictable for patients, staff, and administrations. [32]

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3	332	
4 5	333	Another finding of this systematic review was a significant reduction in the length of hospital
6 7 8	334	stay following patients' examination in a PAC; however, a small number of studies with low
9 10	335	quality were considered. Nevertheless, similar results were found in another systematic
11 12 13	336	review claiming that perioperative systems support hospitals addressing the expected
13 14 15	337	growth in the number and complexity of surgical procedures. [17] When patients are
16 17	338	examined in the PAC and well-prepared with information, consultations, and tests, they
18 19 20	339	need not be hospitalised until the day of surgery. A survey on operated patients showed that
21 22	340	given a choice, 75% do not wish admission to the hospital until the day of operation; a major
23 24 25	341	reason being shorter hospital stay. [41] However, an updated systematic review on the
26 27	342	effectiveness of nurse-led preoperative assessment services for elective surgery found that
28 29 30	343	the included articles demonstrated a reduced length of stay; these studies had low
31 32	344	methodological quality, and therefore, the authors could not conclude that this service leads
33 34 35	345	to reduced length of hospital stay. [18]
36 37	346	
38 39 40	347	The evidence from our systematic review is insufficient to conclude whether patients have
40 41 42	348	reduced anxiety due to PAC. The included studies used different instruments for measuring
43 44 45	349	anxiety levels, and the results could not be pooled. [42]
46 47	350	
48 49 50	351	A major purpose of establishing a PAC in a hospital is to better prepare the patients for the
50 51 52	352	anticipated surgery. Healthcare professionals and policymakers are exploring strategies to
53 54	353	reduce unnecessary investigations without compromising quality and patient safety. [43]
55 56 57	354	Translation of evidence-based interventions into hospital systems can provide instant and
58 59 60	355	substantial benefits to patients care and outcomes. However, existing literature describes

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356	barriers and facilitators to implementation related to the system, staff and the intervention.
357	[40] "Routine preoperative laboratory tests" on relatively healthy patients are not
358	recommended from the American Society of Anesthesiologist (ASA), [16] and the National
359	Institute of Health and Care Excellence (NICE). [44] Instead, they encourage for patient and
360	surgery- specific investigations. This recommendation is not always implemented in hospital
361	protocols or practiced for several reasons. Furthermore, an observational study found that
362	routine pre-operative testing to predict abnormalities found at least one abnormal test
363	results in most of the relatively healthy patients. Only 0.67% of the abnormalities had
364	significant impact on changing the perioperative management. [45] Blitz et al. argued that
365	PACs should focus on early patient engagement strategies, interdisciplinary team
366	communication, detailed perioperative care plans, and patient documentation using
367	electronic health record that should be open for review by the perioperative team to
368	preserve patient information and safety. [15] Furthermore, a previous study pointed out
369	that the risk factors are not only patient-related but also organisation-related, [8] and that
370	some hospitals have perioperative care teams that are better at identifying and relieving
371	perioperative complications. [46,47] This suggests that the value of PACs lies in their ability
372	to improve the quality of the perioperative process by designing a more robust system for
373	preoperative assessment and preparation. [15] A Danish study found that deaths attributed
374	to anaesthesia were associated with inadequate preoperative evaluation. [7] However, the
375	assessment of PAC was significantly associated with reduced mortality following complex
376	orthopaedic surgery in only one study in this systematic review. [35] Retrospective studies
377	have reported similar results, but with different surgeries. [15,48]
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#### 30 Strengths and limitations of the study

1 The review was performed in duplicate or independently by two researchers, and consensus 32 was reached through discussion. However, grey literature, such as government and 33 institutional documents, were excluded and might be a limitation of this study. Since 34 organisation of healthcare systems vary in countries, the type of staff who performed the 35 patient's preoperative assessment was not considered an inclusion criterion. The European Society of Anaesthesiology guidelines recommend that anaesthesiologists complete the 86 37 preoperative assessment, while trained nurses or anaesthesia trainees perform the 88 screening. [8] A preoperative evaluation performed by an internist was associated with increased length of stay and increased postoperative mortality. [49] This systematic review 39 90 results were possibly affected by the heterogeneity in the types of staff performing the preoperative assessment. 1

We exclusively included studies with high internal validity. Therefore, we excluded several 93 4 retrospective studies. Nonetheless, remaining studies' risk of bias was fairly high, and were 95 heterogeneous and meta-analyses were not statistically appropriate. [27] The included 96 studies' designs could not rule out selection bias and confounding, and the strength of the 97 evidence should be assessed cautiously. Many studies did not make adjustments for several 98 confounders, which could be responsible for the observed effects. Several studies lacked 99 descriptions of the methods used and the patients included, lowering transparency. It is not very reassuring that many such studies were unable to deliver more thorough evidence to 00 1 guide practice and should be assessed cautiously. The results are relevant to health care )2 services, focusing on the well-being and safety of the patients.

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3 4	404	Implications for future research and practice
5 6 7	405	This systematic review identified the ambiguity in the PAC interventions offered to the
, 8 9	406	intervention group. In many studies, it was evident that the methods used lacked clarity, and
10 11	407	high-quality research is needed in this field. The studies included in this review did not
12 13 14	408	contain any results of earlier surgical room entry time for patients assessed in PACs, [50,51]
15 16	409	reduced preoperative tests, such as blood tests, in patients before surgery after attending
17 18 19	410	the PAC, similar to previous retrospective studies. [28,52]
20 21	411	Other implications for future research might include the organisation structure of different
22 23 24	412	PACs and their functioning. Additionally, the tests that should be part of assessement at
24 25 26	413	PACs should be investigated. The use of technology, such as streaming services, facilitates
27 28	414	different patient groups and might become crucial for reducing human contact and spread
29 30 31	415	of the virus in context of Covid-19.
32 33	416	CONCLUSION
34 35 36	417	PAC use has reduced the length of stay and cancellation rate at hospitals. However, the
37 38	418	effectiveness of PAC, the major review question, remains unclear, and requires further
39 40 41	419	research. There is a demand for high-quality studies capturing robust data describing the
42 43	420	quality of care and clinical outcomes for patients requiring anaesthesia. This requires
44 45 46	421	increased focus and funding for this specific area of health services research and could,
40 47 48	422	therefore, lead to implementations of PACs in health care services and improve patient
49 50 51	423	safety and perioperative care.
51 52 53	424	Acknowledgement
54 55	425	We thank the librarian Ellen Sejerstedt, University of Agder, Kristiansand, for helping with
56 57 58	426	the search strategy and removal of duplicates.
59 60	427	Competing Interests

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3 ⊿	428	The authors declare that they have no conflict of interest.
4 5		
6 7	429	Data sharing statement
8 9	430	All data relevant to the study are included within the article or have been uploaded within
10 11 12	431	supplemental files.
13 14	432	Funding
15 16 17	433	This research received no specific grant from any funding agency in the public, commercial
18 19	434	or not-for-profit sectors.
20 21 22	435	Author contributions
23 24	436	EWK, MF, AO and TOT: Study design
25 26 27	437	EWK, MF, AO: Search and screening of the articles
27 28 29	438	EWK: Data extraction
30 31	439	MF, AO, RCB and TOT: Control of data extraction
32 33 34	440	EWK, MF, AO and RCB: Quality assessment of the included articles
35 36	441	EWK: Drafting of the manuscript
37 38 30	442	MF, AO, RCB, and TOT: Contribution to and review of the final version of the manuscript
40 41	443	MF, AO, RCB and TOT: Supervised the study
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12	604	Figure 2: The Joanna Briggs Institute Critical appraisal checklist for quasi-experimental
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Figure 2: The Joanna Briggs Institute Critical appraisal checklist for quasiexperimental studies was used for the risk of bias assessment. [30]

# Appendix 1: Search strategies

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<ul> <li>Search update, 4 February 2021</li> <li>A1. Main Databases</li></ul>	5 5 6

# 1. Database Search Strategies

The search mode for CINAHL was Boolean/Phrase. For those searches that are done without search fields - it is automatically searched in the standard fields that CINAHL uses, including words from title, summary, and subject headings. Search 1 and Search 2 are with words in the title (TI in front of the keywords), however this have not been done with the other searches.

Embase and Medline have the same search mode because we are searching for words from title, summary, and subject headings.

## 2. Initial searches, 11 September 2018

### 2.1. Main Databases

Database: Embase, (Ovid)

Results: n=1287

Search:

1 ((pre-operativ\* or preoperativ\* or pre operativ\*) and (assessment\* or measurement\* or evaluat\*)).ti.

2 (clinic\* or unit\* or nurs\* or outpatient\* or ward\* or center\* or centre\*).ti.

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6 ((anaesthe\* or anasthe\* or anesthe\*) adj4 outpatient\* adj4 clinic\*).ti,ab.

7 ((pre-admiss\* or preadmiss\*) adj4 (center\* or centre\*) adj4 (evaluat\* or assessment\* or measurement\*)).ti,ab.

- 8 or/3-7
- 9 limit 8 to yr="1996 -Current"
- 10 limit 9 to (conference abstracts or embase)

#### Database: Medline (Ovid)

Results: n=997

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34		or clinic* or unit* or nurs* or outpatient* or ward* or centre* or center*)) and (surg*
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44 45		TO limit 9 to (conference abstracts of embase)
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4/		Database: Medline (Ovid)
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55		2 (chine of unit of nuis of outpatient of ward of center of centre ).
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58		preanesthe* or pre anesthe*) adj4 (assessment* or measurement* or evaluat* or
59		clinic* or nurs* or unit* or outpatient* or ward* or center* or centre*)) ti.ab.
60		

5 (((pre-admiss\* or preadmiss\*) adj4 (assessment\* or measurement\* or evaluat\* or clinic\* or unit\* or nurs\* or outpatient\* or ward\* or centre\* or center\*)) and (surg\* or anaesthe\* or anesthe\* or preoperativ\* or pre-operativ\* or preanaesthe\* or pre-anaesthe\* or pre-anaesthe\* or pre-anesthe\* or preanesthe\* or pre-anesthe\* or preanesthe\* or pre anesthe\*)).ti,ab.
6 ((anaesthe\* or anasthe\* or anesthe\*) adj4 outpatient\* adj4 clinic\*).ti,ab.
7 ((pre-admiss\* or preadmiss\*) adj4 (center\* or centre\*) adj4 (evaluat\* or assessment\* or measurement\*)).ti,ab. (4)
8 or/3-7

9 limit 8 to yr="1996 -Current"

Database: Cinahl Plus with Full Text (EBSCO host)

Results: n=209

Search:

1 TI ((pre-operativ\* or preoperativ\* or "pre operativ\*") AND (assessment\* or measurement\* or evaluat\*))

2 TI (clinic\* or unit\* or nurs\* or outpatient\* or ward\* or center\* or centre\*)
3 S1 AND S2

4 (preanaesthe\* or pre-anaesthe\* or "pre anaesthe\*" or pre-anesthe\* or preanesthe\* or "pre anesthe\*") N3 (assessment\* or measurement\* or evaluat\* or clinic\* or nurs\* or unit\* or outpatient\* or ward\* or centre\* or center\*)

5 ((pre-admiss\* or preadmiss\*) N3 (assessment\* or measurement\* or evaluat\* or clinic\* or unit\* or nurs\* or outpatient\* or ward\* or centre\* or center\*)) AND (surg\* or anaesthe\* or anesthe\* or preoperativ\* or "pre operativ\*" pre-operativ\* or preanaesthe\* or pre-anaesthe\* or pre-anesthe\* or preanesthe\* or "pre anesthe\*") 6 (anaesthe\* or anasthe\* or anesthe\*) N3 outpatient\* N3 clinic\* 30

7 (pre-admiss\* or preadmiss\*) N3 (center\* or centre\*) N3 (evaluat\* or

assessment\* or measurement\*)

8 S3 OR S4 OR S5 OR S6 OR S7

9 S3 OR S4 OR S5 OR S6 OR S7 Limiters - Published Date: 19960101-; Exclude MEDLINE records

# Reporting checklist for systematic review and meta-analysis.

Based on the PRISMA guidelines.

# Instructions to authors

Complete this checklist by entering the page numbers from your manuscript where readers will find each of the items listed below.

Your article may not currently address all the items on the checklist. Please modify your text to include the missing information. If you are certain that an item does not apply, please write "n/a" and provide a short explanation.

Upload your completed checklist as an extra file when you submit to a journal.

In your methods section, say that you used the PRISMAreporting guidelines, and cite them as:

Moher D, Liberati A, Tetzlaff J, Altman DG, The PRISMA Group. Preferred Reporting Items for Systematic Reviews and Meta-Analyses: The PRISMA Statement

		Page
	Reporting Item	Number
Title	4	
<u>#*</u>	Identify the report as a systematic review, meta-analysis, or both.	1
Abstract		
Structured #2 summary	Provide a structured summary including, as applicable: background; objectives; data sources; study eligibility criteria, participants, and interventions; study appraisal and synthesis methods; results; limitations; conclusions and implications of key findings; systematic review registration number	2
Introduction		
Rationale #3	<u>B</u> Describe the rationale for the review in the context of what is already known.	3-5
F	or peer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml	

1 2 3 4 5	Objectives	<u>#4</u>	Provide an explicit statement of questions being addressed with reference to participants, interventions, comparisons, outcomes, and study design (PICOS).	5
6 7	Methods			
8 9 10 11 12	Protocol and registration	<u>#5</u>	Indicate if a review protocol exists, if and where it can be accessed (e.g., Web address) and, if available, provide registration information including the registration number.	5
13 14 15 16 17 18 19	Eligibility criteria	<u>#6</u>	Specify study characteristics (e.g., PICOS, length of follow- up) and report characteristics (e.g., years considered, language, publication status) used as criteria for eligibility, giving rational	6
20 21 22 23 24 25 26	Information <u>#7</u> sources		Describe all information sources in the search (e.g., databases with dates of coverage, contact with study authors to identify additional studies) and date last searched.	5,6
27 28 29 30 31	Search	<u>#8</u>	Present full electronic search strategy for at least one database, including any limits used, such that it could be repeated.	5,6 + APPENDIX 1
32 33 34 35 36 37 38	Study selection	<u>#9</u>	State the process for selecting studies (i.e., for screening, for determining eligibility, for inclusion in the systematic review, and, if applicable, for inclusion in the meta-analysis).	6
<ol> <li>39</li> <li>40</li> <li>41</li> <li>42</li> <li>43</li> <li>44</li> <li>45</li> </ol>	Data collection process	<u>#10</u>	Describe the method of data extraction from reports (e.g., piloted forms, independently by two reviewers) and any processes for obtaining and confirming data from investigators.	6-7
46 47 48 49 50 51	Data items	<u>#11</u>	List and define all variables for which data were sought (e.g., PICOS, funding sources), and any assumptions and simplifications made.	5-7
52 53 54 55 56 57 58 59 60	Risk of bias in individual studies	<u>#12</u> For p	Describe methods used for assessing risk of bias in individual studies (including specification of whether this was done at the study or outcome level, or both), and how this information is to be used in any data synthesis.	6-7
00		p		

Page 37 of 37

### BMJ Open

Summary measures	<u>#13</u>	State the principal summary measures (e.g., risk ratio, difference in means).	N/A
Planned methods of analyis	<u>#14</u>	Describe the methods of handling data and combining results of studies, if done, including measures of consistency (e.g., I2) for each meta-analysis.	7
Risk of bias across studies	<u>#15</u>	Specify any assessment of risk of bias that may affect the cumulative evidence (e.g., publication bias, selective reporting within studies).	12
Additional analyses	<u>#16</u>	Describe methods of additional analyses (e.g., sensitivity or subgroup analyses, meta-regression), if done, indicating which were pre-specified.	N/A
Results			
Study selection	<u>#17</u>	Give numbers of studies screened, assessed for eligibility, and included in the review, with reasons for exclusions at each stage, ideally with a <u>flow diagram</u> .	8
Study characteristics	<u>#18</u>	For each study, present characteristics for which data were extracted (e.g., study size, PICOS, follow-up period) and provide the citation.	N/A
Risk of bias within studies	<u>#19</u>	Present data on risk of bias of each study and, if available, any outcome-level assessment (see Item 12).	12 + Figure
Results of individual studies	<u>#20</u>	For all outcomes considered (benefits and harms), present, for each study: (a) simple summary data for each intervention group and (b) effect estimates and confidence intervals, ideally with a forest plot.	12-16
Synthesis of results	<u>#21</u>	Present the main results of the review. If meta-analyses are done, include for each, confidence intervals and measures of consistency.	12-16
Risk of bias	<u>#22</u>	Present results of any assessment of risk of bias across	N/A
across studies		studies (see Item 15).	
Additional	<u>#23</u>	Give results of additional analyses, if done (e.g., sensitivity	N/A
analysis		or subgroup analyses, meta-regression [see Item 16]).	
Discussion			
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1 2 3 4 5 6	Summary of Evidence	<u>#24</u>	Summarize the main findings, including the strength of evidence for each main outcome; consider their relevance to key groups (e.g., health care providers, users, and policy makers	16-19
7 8 9 10 11 12	Limitations	<u>#25</u>	Discuss limitations at study and outcome level (e.g., risk of bias), and at review level (e.g., incomplete retrieval of identified research, reporting bias).	19
13 14 15 16	Conclusions	<u>#26</u>	Provide a general interpretation of the results in the context of other evidence, and implications for future research.	20
17 18	Funding			
19 20 21 22 23	Funding	<u>#27</u>	Describe sources of funding or other support (e.g., supply of data) for the systematic review; role of funders for the systematic review.	21
$\begin{array}{c} 25\\ 26\\ 27\\ 28\\ 29\\ 30\\ 31\\ 32\\ 33\\ 34\\ 35\\ 36\\ 37\\ 38\\ 39\\ 40\\ 41\\ 42\\ 43\\ 44\\ 45\\ 46\\ 47\\ 48\\ 49\\ 50\\ 51\\ 52\\ 53\\ 54\\ 55\\ 56\end{array}$	License CC-BY. T made by the EQL	This che	Acklist can be completed online using https://www.goodreports Network in collaboration with Penelope.ai	<u>.org/</u> , a tool
58 59 60		For p	eer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml	

# **BMJ Open**

# Effectiveness of pre-anaesthetic assessment clinic: A systematic review of randomised and non-randomised prospective controlled studies

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Date Submitted by the Author:	16-Mar-2022
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<b>Primary Subject Heading</b> :	Anaesthesia
Secondary Subject Heading:	Surgery, Nursing, Health services research
Keywords:	Adult anaesthesia < ANAESTHETICS, Health & safety < HEALTH SERVICES ADMINISTRATION & MANAGEMENT, Quality in health care < HEALTH SERVICES ADMINISTRATION & MANAGEMENT, Adult surgery < SURGERY
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2 3 4	1	Effectiveness of pre-anaesthetic assessment clinic: A systematic review of randomised and
5 6 7	2	non-randomised prospective controlled studies
7 8 9	3	
10 11	4	Eirunn Wallevik Kristoffersen <sup>1,2</sup> *, Anne Opsal <sup>1</sup> , Tor Oddbjørn Tveit <sup>1,2,3</sup> , Rigmor C Berg <sup>4,5</sup> ,
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47 48	19	E-mail: eirunn.w.kristoffersen@uia.no
49 50	20	
51 52 53	21	Word count: 4156
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3 4	24	ABSTRACT
5	25	Objectives: The aim of this systematic review was to examine the effectiveness of pre-
7	26	anaesthesia assessment clinics (PACs) in improving the quality and safety of perioperative
8 9	27	patient care.
10 11	28	Design: Systematic review.
12 13	29	Data sources: The electronic databases CINAHL Plus with Full Text (EBSCOhost), Medline,
14 15	30	and Embase (OvidSP) were systematically searched on 11 September 2018
16	31	and updated on 3 February 2020 and 4 February 2021.
17 18 19 20 21 22 23 24 25 26 27 28 29 20	32	Eligibility criteria: The inclusion criteria for this study were studies published in English or
	33	Scandinavian language and scientific original research that included randomised or non-
	34	randomised prospective controlled studies. Additionally, studies that reported the outcomes
	35	from a PAC consultation with the patient present were included.
	36	Data extraction and synthesis: Titles, abstracts, and full texts were screened by a team of
	37	three authors. Risk of bias was assessed using the Joanna Briggs Institute critical appraisal
	38	checklist for quasi-experimental studies. Data extraction was performed by one author and
30 31	39	checked by four other authors. Results were synthesised narratively owing to the
32 33	40	heterogeneity of the included studies.
34 35	41	Results: Seven prospective controlled studies on the effectiveness of PACs, were included.
36 37	42	Three studies reported a significant reduction in the length of hospital stay and two studies
38	43	reported a significant reduction in cancellation of surgery for medical reasons when patients
39 40	44	were seen in the PAC. In addition, the included studies presented mixed results regarding
41 42	45	anxiety in patients. Most studies had a high risk of bias.
43 44	46	Conclusion: This systematic review demonstrated a reduction in the length of hospital stay
45 46	47	and cancellation of surgery when the patients had been assessed in the PAC. There is a need
47	48	for high-quality prospective studies to gain a deeper understanding of the effectiveness of
49	49	PACs.
50 51	50	PROSPERO registration number: CRD42019137724
52 53	51	
54 55	52	Keywords: pre-anaesthetic assessment clinic, preoperative care, quality, safety, systematic
56 57	53	review
58 50	54	
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2 3	56	
4	57	
6	58	
7	59	Change at he and light to the of this study.
8 9	60	Strengths and limitations of this study
10 11	61	<ul> <li>An extensive database search was conducted with no limitations on outcomes and</li> </ul>
12 12	62	the type of pre-anaesthetic assessment clinic.
13 14 15	63	Only randomised or non-randomised prospective controlled studies were included.
15 16	64	• The Joanna Briggs Institute Critical appraisal checklist for quasi-experimental studies
17 18	65	was used.
19 20	66	• The included studies were heterogeneous and had a high risk of bias, which is a
21 22	67	major limitation of this review.
23	68	
24 25 26	69	INTRODUCTION
27 28	70	Anaesthesia is crucial in surgery. However, it may activate physiological changes that
29 30 31	71	increase morbidity and mortality,[1] depending on the patients' preoperative health status
32 33	72	and age.[2] Hospitals treat patients with complex, comorbid healthcare problems,
34 35 36	73	undergoing progressively extensive surgeries and interventions.[3,4] To ensure the quality
37 38	74	and safety of anaesthesia and surgery, precise knowledge of the clinical characteristics of
39 40 41	75	patients is critical to the perioperative management.[2] Over the past 50 years,
42 43	76	perioperative mortality, including anaesthesia-related mortality, has declined, which is
44 45	77	significant in developed countries,[1,5] mainly because of new anaesthetics, improved
40 47 48	78	monitoring equipment and training, availability of recovery rooms, and improved airway
49 50	79	management.[4] However, a previous review found higher rates of morbidity and mortality
51 52 53	80	in non-operating room anaesthesia, which was attributed to limited preoperative
54 55	81	evaluation.[6] A retrospective study found significant associations between perioperative
56 57 58	82	mortality and age < 1 year or > 65 years, American Society of Anesthesiologists Physical
59 60	83	Status Classification System (ASA), emergency case status, and operative start time after

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84 6:00 PM.[7] This might indicate that risk factors are both patient- and surgery-related and 85 may be linked to organisational structures.[8]

86 Currently, an increasing number of pre-anaesthesia assessment clinics (PACs) support 87 hospitals internationally in handling the rising number of patients and complexity of surgical 88 procedures.[9] The design of PACs differs critically based on location, organisational 89 structure, timing, and patient groups. They primarily function as a service unit for surgeons, 90 patients, and the anaesthetic team.[10] The PAC consultation, by the anaesthesiologist, 91 anaesthesia nurse, or both, is a globally recognised evaluation method and optimises the 92 patients' medical condition prior to surgery and anaesthesia.[2] Thus, it is essential for 93 secure anaesthetic practice since it detects anaesthesia-related risk factors and high-risk 94 patients, improves patient outcomes, prepares the patient physically and psychologically for 95 anaesthesia, and ensures the patient's most favourable condition for surgery and 96 anaesthesia.[11-13] This is primarily performed by interviewing and examining the patient; 97 reviewing previous medical, surgical and anaesthesia issues; evaluating current medication; 98 and obtaining and reviewing preoperative tests.[10] PACs also allow increased 99 communication between healthcare providers and coordination with postoperative .00 care.[14,15] Because of well-prepared patients and staff, several researchers have indicated .01 that with PAC, the number of surgical cancellations, length of hospital stay, laboratory tests, .02 and mortality rate have reduced. [7,16,17] Others assert that patients feel less anxious .03 regarding the subsequent anaesthetic and surgical processes and are highly satisfied with .04 the service with PAC consultations.[16,18,19]

.05 As Turunen et al. stated, research on PACs regarding costs, financial savings, the impact on .06 patient safety and quality of care, accuracy of the number of operative patients, and effect

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107	on preoperative nursing levels, is scarce.[20] Survey results indicate that anaesthesiologists
108	perceive day of surgery delays due to missing information as common, even with PAC
109	consultations.[21] This systematic review aimed to examine the effectiveness of PACs in
110	improving the quality and safety of perioperative patient care. Further, we aimed to
111	determine the gaps in existing knowledge for future research.
112	METHODS
113	Our systematic review followed the guidelines in the Cochrane Handbook for Systematic
114	Reviews of Interventions[22] and was reported in accordance with the Preferred Reporting
115	Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines.[23] The protocol was
116	registered in PROSPERO: CRD42019137724.[24]
117	The two review questions were:
118	1. Is PAC effective in improving patient satisfaction and safety, while reducing anxiety?
119	2. Is PAC effective in reducing cancellation rate and cost of surgery, and improving the
120	efficiency of perioperative patient care?
121	Search strategies
122	We performed a scoping search in different databases to identify the key terms.[25,26] The
123	final search was planned and conducted in close collaboration with the university librarian.
124	On 11 September 2018 we searched CINAHL Plus with Full Text (EBSCOhost), Medline, and
125	Embase (OvidSP) databases, which were updated on 3 February 2020 and 4 February 2021.
126	Considering the lack of subject headings (e.g., MeSH) for PAC, we combined text words, such
127	as 'pre-anaesthesia', 'nurse', 'surgery', 'anaesthesia', 'preoperative', 'assessment',
128	'measurement', 'evaluate', 'preadmission', 'centre', 'clinic', 'ward', 'unit', and 'outpatient'.
129	The searches are detailed in Appendix 1. The search mode in CINAHL was Boolean/Phase,

Page 7 of 40

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3 4	130	which supports Boolean searching or exact phrase searching. For comprehensiveness, we
5	101	used both the truncetion and every insity encycles . We livelted the second to 1000 the year
6 7	131	used both the truncation and proximity operators. We limited the search to 1996, the year
7 8	132	one of the first known articles in this area was published. [27] Complementary methods to
9	102	one of the mot known at these in this area was published.[27] complementary methods to
10 11	133	identify studies included following up on citations via Scopus, scanning the reference lists of
12		
13	134	relevant papers and included articles, and checking for relevant studies in clinical trials.[25]
14 15		
16	135	Eligibility criteria
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18 19	136	The main inclusion criterion was that the study, using empirical quantitative methods,
20	127	addressed the effectiveness of PACs. Specific eligibility criteria were: (a) published in English
21 22	137	addressed the effectiveness of FACs. Specific enginity criteria were. (a) published in English
22	138	or Scandinavian language, (b) scientific publication of original research, (c) reported the
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25 26	139	outcomes of PAC, (d) PAC consultation with the patient present, (e) randomised or non-
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28	140	randomised prospective controlled studies, and (f) newly established PAC. The following
29 30		
31	141	were excluded: (a) editorials, discussions, and conference abstracts, (b) reviews, (c)
32	142	instrument testing (d) studies on children and (e) retrospective studies
33 34	172	instrument testing, (a) studies on enharch, and (e) retrospective studies.
35	143	Study selection
36 37		
38	144	All references identified in the search were transferred to EndNoteX9, where the duplicates
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40 41	145	were removed. Subsequently, all unique references were transferred to the Covidence
42	116	ccrooning tool [29] Study aligibility was accortained independently, first at the title and
43	140	screening tool.[26] Study enginity was ascertained independently, inst at the title and
44 45	147	abstract level, and subsequently at the full text level. Three of the authors screened all the
46		
47 48	148	articles (EWK, AO, MF). Inclusion was determined by consensus, and disagreements were
49		
50	149	resolved by consulting two other authors (RCB and TOT).
51 52	450	
53	150	Quality assessment
54	151	Rick of higs in studies were assessed using design-specific checklists. Given the
55 56	151	hisk of bids in studies were disessed danig design specific encekists. Given the
57	152	methodological similarity of the studies, only the Joanna Briggs Institute Critical appraisal
58 50		
60	153	checklist for quasi-experimental studies was used.[29] Author EWK performed the risk of

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3 4	154	bias assessment, and RCB confirmed its accuracy. Disagreements were resolved through
5 6 7	155	discussion with MF and AO. Each of the nine checklist questions was answered no, yes,
8 9	156	unclear, or not applicable.
10 11 12	157	Data extraction and analysis
12 13 14	158	Author EWK extracted data from each study onto a pre-designed Excel spreadsheet. All the
15 16	159	authors confirmed the accuracy, consistency, and completeness of the extracted data that
17 18 19	160	included publication details; study design; setting; and characteristics of the patients,
20 21	161	interventions, comparisons, and outcome (PICO). We requested information on the missing
22 23 24	162	data; however, received no response from the authors. If the PICO elements had been
25 26	163	sufficiently similar and statistical data were available, we had intended to conduct a meta-
27 28 20	164	analysis. However, the extracted data revealed substantial heterogeneity. Therefore, we
29 30 31	165	performed a narrative synthesis. We described the findings in text, stratified by outcome,
32 33	166	with descriptions of the effects of interventions for each study, classification of the effect
34 35 36	167	direction, and we looked across contributing studies to develop a summary of findings for
37 38	168	each outcome.[22]
39 40 41	169	Patient and Public Involvement
42 43	170	Patients and/or the public were not involved in the design, conduct, reporting, or
44 45 46	171	dissemination plans of this research. However, the project was initiated by health
40 47 48	172	professionals.
49 50	173	
52 53	174	RESULTS
54 55	175	Figure 1. provides the details of the study selection process. A total of 2,981 records were
56 57 58 59 60	176	identified in the final search (2021). After removing duplicates, we screened 2,058 records

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The seven studies are listed in Table 1. All seven studies were in English and were published

report data collection information).[30] Based on the inclusion criteria, all were prospective

between 2000 and 2017, with data collected between 1997 and 2015 (one study did not

controlled studies; however, no RCTs were found. There was one controlled before-after

implementation, without baseline assessments. There were three two-group non-parallel

after-only studies, [32-34] and three two-group parallel after-only studies; [30] one had a

matched control group[35] and one had three follow-up assessments of one arm.[36] One

study had only cancellation rate as prospective data.[32] The studies included 77,411

ata.

study.[31] The remaining six studies had control groups; assessments followed PAC

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patients.

based on the title and abstract; 179 records passed the full-text screening. After applying theinclusion criteria, seven studies were selected for the final analysis.

**Overall characteristics of the studies** 

## 190 Table 1: Description of the included studies

Author, Year, Country	Study design	Sampling time	Population	Intervention	Comparison	Outcomes
Farasatkish,2009[33] Iran	Two-group after study	May 2007 through August 2007	N=1716, open- heart surgery, ASA class III-IV	Pre-anaesthesia consultation clinic (3-10 days before surgery)	Usual care (within 24 h of surgery)	Cancellations
Kamal,2011[34] England	Two-group	April 2005 through April 2009	N=1445, complex elective orthopaedic surgery, ASA class III-IV	Preoperative anaesthetic assessment clinic (timing not stated)	Usual care (day of surgery)	Admissions, length of stay, mortality, cost
Kamau,2017[31] Kenya	CBA	August 2000, April 2001, November 2001	N=51, elective non-cardiac surgery, ASA class not stated	Pre-anaesthesia clinic consultation (≥48 h before surgery)	Usual care (day before surgery)	Anxiety (STAI score)
Klopfenstein,2000[30] Switzerland	Two-group after study (parallel)	No data	N=40, elective endoscopic urological surgery, ASA class I-III	Pre-anaesthetic consultation (1-2 weeks before surgery)	Usual care (the evening before surgery)	Anxiety (MAACL, VAS)
Lee,2012[35] China	Two-group after study (parallel)	March 2007 through November 2009	N=352, elective surgery, ASA class I-IV	Anaesthesia consultation clinic (≤3 months before surgery)	Usual care (the evening before surgery)	Quality of recovery score, cost, cancellations, length of stay, satisfaction, anxiety (VAS), willingness to pay (WTP)
Mendes,2005[36] Brazil	Two-group after study (parallel)	April 2007 through June 2007	N=52254, surgery, ASA class not stated	Preoperative outpatient evaluation clinic (timing not stated)	Usual care (timing not stated)	Cancellations, length of stay

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2 3 4 5 6 7 8	191	van Klei,2002[32] The Netherlands	Two-group after study	November 2012	N=21553, elective surgery, ASA class not stated	Preoperative outpatient evaluation clinic (average 3 weeks before surgery)	Usual care (day before surgery)	Cancellations
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10 11 12 13 14 15	192 193	ASA: American Societ Inventory; VAS: visual	y of Anesthesiolo l analogue scale;	ogy; CBA: controlle WTP: willingness	ed before-after; M to pay	AACL: Multiple Affect Adj	ective Check List; S <sup>-</sup>	Al: State-Trait Anxiety
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194	Of the 77,411 patients in the studies, 9,626 and 15,531 patients were in the intervention and
195	control groups, respectively. One study did not specify the number of patients in the
196	intervention and control groups, but only the total number of surgeries performed.[36] Five
197	studies reported data for sex, showing that 51% of the patients were women and 49% were
198	men (12,129 vs. 11,583).[30-33,35] There were more women than men in both the
199	intervention (4,345 vs. 4,134) and control groups (7,784 vs. 7,449). Five studies reported
200	data for age showing that all patients were over 20 years of age[30-33,35] and four studies
201	had grouped within the American Society of Anesthesiology (ASA) category.[30,33-35]
202	
203	The patients were scheduled to undergo a variety of surgeries, including
204	orthopaedic,[31,32,34,35] urology,[30-32,35] general,[31,32,35] heart,[33]
205	gynaecology/obstetrics,[31,32,35] vascular surgery,[32] ophthalmology,[32]
206	maxillofacial/dental surgery,[31,32] and neurological surgery,[32] while one did not specify
207	the type.[36] In five studies, the type of anaesthesia was not specified,[31-34,36] and two
208	studies reported patients for general and/or regional supplement.[30,35]
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210	The patients included had previous anaesthetic experience in one study,[30] previous and no
211	previous anaesthetic experience in another, [31] and five studies did not report this data. [32-
212	36] Limited background characteristics of the patients were reported in two studies.[34,36]
213	One stated that the patients included had ASA 3 or 4 and a body mass index >40; however,
214	no ASA number, sex, or age was reported.[34] Mendes et al. did not report any background
215	characteristics of the included patients.[36]
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217	Considering the intervention, the PACs in all studies comprised an outpatient service
218	whereby patients were examined for medical conditions important for anaesthesia and
219	informed regarding expectations on the day of surgery. Nevertheless, the terminology used
220	for PACs varied, as they served different surgical specialities and conducted pre-anaesthesia
221	consultation from $\geq$ 48 h to $\leq$ 3 months before the surgery. The settings included a university
222	hospital (n=3),[31,35,36] teaching hospital (n=1),[32] medical centre (n=1),[33] and general
223	hospital (n=1);[34] one study did not specify the context.[30] The staff conducting the pre-
224	anaesthetic consultation also varied: in five studies, it was the anaesthesiologists,[30,32,34-
225	36] in the other studies, it was (also) the orthopaedic senior house officer,[34] consultant or
226	resident,[31] or physician.[33] In three studies, nurses were part of the team.[32,34,35] The
227	comparison in all studies was usual care, which generally involved a preoperative
228	anaesthetic evaluation of the admitted patients the day before the surgery.
229	Description of risk of bias in the studies
229 230	Description of risk of bias in the studies Figure 2. shows the results of the risk of bias assessment. In all seven included studies, the
229 230 231	Description of risk of bias in the studies Figure 2. shows the results of the risk of bias assessment. In all seven included studies, the cause and effect were clear. Most of the studies measured outcomes similarly and used
229 230 231 232	Description of risk of bias in the studies         Figure 2. shows the results of the risk of bias assessment. In all seven included studies, the         cause and effect were clear. Most of the studies measured outcomes similarly and used         appropriate statistical analyses. Several studies had limitations of follow-up and similarity in
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229 230 231 232 233 234 235 236 237 238	Description of risk of bias in the studiesFigure 2. shows the results of the risk of bias assessment. In all seven included studies, thecause and effect were clear. Most of the studies measured outcomes similarly and usedappropriate statistical analyses. Several studies had limitations of follow-up and similarity incare and participants. None of the patients had multiple pre-and post-measurements.Outcomes of the included studies are each described separately below.SatisfactionOne study reported satisfaction as an outcome.[35] The summarised patient satisfactionwith the anaesthetic consultation score out of 100 showed that the patients in the PAC
229 230 231 232 233 234 235 236 237 238 239	Description of risk of bias in the studies         Figure 2. shows the results of the risk of bias assessment. In all seven included studies, the         cause and effect were clear. Most of the studies measured outcomes similarly and used         appropriate statistical analyses. Several studies had limitations of follow-up and similarity in         care and participants. None of the patients had multiple pre-and post-measurements.         Outcomes of the included studies         The outcomes of the included studies are each described separately below.         Satisfaction         One study reported satisfaction as an outcome.[35] The summarised patient satisfaction         with the anaesthetic consultation score out of 100 showed that the patients in the PAC         group were more satisfied (mean difference, 2.10%; 95% confidence interval [CI], 0.51–

in mean patient satisfaction with perioperative anaesthesia care score after surgery (mean difference 0.01%, p=0.94).[35] The quality of recovery (QoR) measure referred to the patients' quality of recovery score.[37] The mean QoR score (range, 0–18) following anaesthesia on the first day after surgery was similar between the intervention (13.17±2.73) and control (13.31±2.65) groups (*p*=0.67).[35] Anxiety Three studies reported anxiety. [30,31,35] Two studies reported the visual analogue scale (VAS), one rated from zero (no anxiety) to ten (very high anxiety),[30] another used a 100-mm horizontal line with 'not anxious at all' to 'extremely anxious'.[35] In one study, the median VAS anxiety score was 3 (0–5) in the intervention group and 5 (2–8) in the control group (p=0.0038).[30] In another study, there were no significant differences between the control and intervention groups for levels of anxiety (VAS), surgery (26 vs. 25, respectively, p=0.12), and anaesthesia (20 vs. 19, respectively, p=0.60).[35] The median Multiple Affect Adjective Check List (MAACL) score, with possible range of scores from 0 to 21 (higher scores indicating greater levels of anxiety), was 3 (0–9) in the intervention group and 6.5 (2–12) in the control group (p=0.0053).[30] The differences in the State-Trait Anxiety Index (STAI) score, which comprised 40 questions rated on a four-point Likert scale, was 1.51 (95% CI: 1.02–2.02%, p=0.0051).[31] The results on anxiety in these two studies were significant. However, Kamau et al. found no differences on examining anxiety and the influences of sex, duration of hospital stay, and prior anaesthesia experience.[31] Mortality One study reported the mortality rates.[34] Patients attending the High Dependency Unit (HDU), Intensive Care Unit (ICU), and Post-anaesthesia Care Unit (PACU) following complex 

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264	orthopaedic surgery had a significant reduction in mortality rate after being assessed at the
265	PAC, from 18 (6.1%) of 298 patients to 14 (1.2%) of 1147 patients ( <i>p</i> =0.001).[34]
266	Cancellation rate
267	Four studies reported reduced cancellation rates following the establishment of a
268	PAC.[32,33,35,36] One of the included studies had 316 (2.0%) cancellations for medical
269	reasons before the introduction of PAC, and 79 (0.9%) after, with a difference of 1.02% (95%
270	CI, 0.31–1.31%). After adjustment, the odds ratio was 0.7 (95% CI, 0.5–0.9%).[32] The overall
271	cancellation of surgery reduced from 1027 (6.3%) to 393 (4.6%) following PAC introduction,
272	with a difference of 0.9% (95% Cl, 0.3–1.0%).[32] Mendes <i>et al</i> .[36] found a decrease in
273	overall cancellations from year 1 (39.3%) to year 4 (15.9%), $p \le 0.05$ . In the first year of their
274	study, there were 469 cancellations per 10,639 surgeries performed. The following year, a
275	considerable increase above the baseline in the intervention group was observed, followed
276	by a progressive decrease in the last year with 391 cancellations per 10,397 surgeries
277	performed.[36] Farasatkish et al. reported that of the 1,716 patients studied, a mean of 15.1
278	% cancelled in the two groups. The cancellation rates in the control and intervention groups
279	were 16.8% (146 [number of cancellations]/866 [number of surgeries]) and 13.29%
280	(113/850) (p=0.046), respectively. The most common reason for cancellation was
281	incomplete medical work-up; 51/146 (35%) in the control group and 32/113 (28%) in the
282	intervention group (p=0.03).[33] Lee <i>et al.</i> found similar rates for surgery being cancelled on
283	the scheduled date for the intervention group compared with the control group (2.3% vs.
284	3.4%, <i>p</i> =0.75).[35]
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286	Costs and willingness to pay

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287	Two studies reported the costs.[34,35] One study reported a total saving of $\pm$ 486.62 per
288	patient after establishing a PAC.[34] Another reported a significantly lower preoperative cost
289	per patient in the intervention group compared with that of the control group (mean
290	difference <i>,</i> \$ 463; 95% Cl, -\$648 to -\$278 per patient, <i>p</i> <0.01).[35] However, the mean
291	difference in the total perioperative treatment cost was not significant, even after adjusting
292	for cancellation on the day of surgery costs.[35] Compared to the control group, the
293	willingness to pay (WTP) among the intervention group patients was significantly more than
294	the median WTP (US \$13) for a clinic consultation at the PAC.[35]
295	Length of stay
296	The length of stay was reported in three studies.[34-36] Mendes et al.[36] found a significant
297	decrease in mean hospital stay of patients from 6.2 to 5.0 days (p $\leq$ 0.001) during the 4 years
298	of this study. Kamal <i>et al.</i> [34] found a significant reduction in the length of stay in the HDU
299	from 2.1 days to 1.6 days ( $p$ =0.01), and in the ICU from 2.3 days to 1.9 days ( $p$ =0.01). In the
300	last study, no significant changes were found in the median duration of postoperative stay
301	between the intervention and control groups.[35]
302	Organisation planning and efficiency
303	Organisation planning and efficiency have been reported in two studies.[34,36] One study
304	found statistically significant changes in the reduction of unplanned admissions to the PACU
305	(65/298 [22%], 111/1147 [10%], p=0.001), ICU (4/298 [1.3%], 4/1147 [0.4%], p=0.01), and
306	HDU (4/298 [1.34%], 20/1147 [1.7%], <i>p</i> =0.01) after implementing a PAC.[34] The planned
307	admissions in the ICU (4/298 [1.3%], 18/1147 [1.6%], <i>p</i> =0.01), and HDU (14/298 [4.7%],
308	85/1147 [7.4%], <i>p</i> =0.1) increased after implementing a PAC.[34] The number of PAC
309	evaluations increased from 14,704 (year 1) to 413,990 (year 4) ( $p \le 0.001$ ).[36] The number

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3 4	310	of outpatient procedures increased from 2,170 (year 1) to 1,943 (year 4) ( $p \le 0.001$ ) and the
5 6 7	311	inpatient procedures decreased from 9,556 (year 1) to 8,449 (year 4) ( $p \le 0.001$ ).[36]
7 8 9 10	312 313	DISCUSSION
11 12 13	314	This systematic review summarises the effectiveness of PACs in improving quality and safety
14 15 16 17	315	of pre-anaesthetic patient care in general hospitals and determines the gaps in existing
	316	knowledge for future research. Herein, we present the main results of the review and infer
18 19 20	317	the implications for research and practice.
21 22	318	
23 24 25 26 27 28 29 30 31 32 33 34 35 36 37	319	Seven studies met the inclusion criteria, and the main findings were reduction in the length
	320	of stay and surgery cancellation rate in hospitals. However, the studies were of low quality,
	321	making it difficult to draw any conclusion. The evidence from our systematic review is
	322	insufficient to conclude whether patients have reduced anxiety because of PAC. This is
	323	because the included studies used different instruments for measuring anxiety levels, and
	324	the results could not be pooled.[38]
38 39	325	
40 41 42	326	A major purpose of establishing a PAC in a hospital is to better prepare the patients for the
43 44	327	anticipated surgery. Healthcare professionals and policymakers are exploring strategies to
45 46 47	328	reduce unnecessary investigations without compromising quality of care and patient
47 48 49	329	safety.[39] Transition of evidence-based interventions to the hospital systems can provide
50 51	330	substantial benefits to patient care.[40] According to the ASA and the National Institute of
52 53 54	331	Health and Care Excellence, routine preoperative laboratory tests are not recommended for
55 56	332	relatively healthy patients. Instead, they encourage patient and surgery- specific
57 58 59	333	investigations.[15,40] This recommendation is not always implemented in hospital protocols
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2 3	334	or practice. An observational study showed that routine preoperative testing to predict
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6 7	335	abnormalities found at least one abnormal test result in most of the relatively healthy
8 9	336	patients. Only 0.67% of the abnormalities had a significant impact on changing the
10 11 12	337	perioperative management.[41] Blitz et al. argued that PACs should focus on early patient
12 13 14 15 16	338	engagement strategies, interdisciplinary team communication, detailed perioperative care
	339	plans, and patient documentation using electronic health record, which should be open for
17 18 19	340	review by the perioperative team.[14] Furthermore, a previous study mentioned that the
20 21 22	341	risk factors are not only patient-related but also organisation-related,[7] and that some
22 23 24	342	hospitals have perioperative care teams that are better at identifying and relieving
25 26	343	perioperative complications.[42,43] Thus, the value of PACs lies in their ability to improve
27 28 29 30 31 32 33 34 35 36	344	the quality of the perioperative process by designing a more robust system for preoperative
	345	assessment and preparation.[14] A narrative review found higher rates of morbidity and
	346	mortality in non-operating room anaesthesia, and one of the main reasons was limited
	347	preoperative evaluation.[6] In this systematic review, the assessment of PAC was
37 38 39	348	significantly associated with reduced mortality following complex orthopaedic surgery in
40 41	349	only one study.[34] Retrospective studies have reported similar results, but with different
42 43 44	350	surgeries.[14,44]
45 46	351	
47 48 40	352	Cancellation on the day of surgery has undesirable effects on both the patients and the
49 50 51	353	hospital system.[13] Late patient-related cancellations can totally or partially be
52 53	354	prevented,[45] if addressed during preoperative evaluations.[16] This has been confirmed by
54 55 56	355	only three studies in this systematic review that found a reduction in surgery cancellation
57 58	356	after implementing a PAC.[32,33,36] However, Lee et al. found no significant changes
59 60	357	between the intervention and control groups.[35] Mendes et al. found that the number of

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58 cancellations for medical reasons after PAC implementation decreased in the first year of 59 implementation. In the second and third years, they were high before the number dropped to below baseline.[36] These conflicting findings indicate that hospitals operate in specific 60 51 contexts, with unique populations, processes, and microsystems, encountering unique 52 obstacles, making implementation difficult. Patient-focused interventions should consider 63 barriers, facilitators, and interrelationships between systems, staff, and interventions to 54 increase the likelihood of sustainable success.[46] Additionally, Kamal et al. indicated that 65 PACs lead to more planned admissions to the ICU, HDU, and PACU, which is more predictable for patients, staff, and administration.[34] 66 67 68 Another finding of this review was a significant reduction in the length of hospital stay 59 following patients' examination in a PAC; however, a small number of studies with low 70 quality were considered. Nevertheless, similar results were found in another systematic 71 review claiming that perioperative systems support the hospitals by addressing the expected 72 growth in the number and complexity of surgical procedures.[16] When patients are 73 examined in the PAC and well-prepared with information, consultations, and tests, they 74 need not be hospitalised until the day of surgery. A survey on operated patients showed that 75 given a choice, 75% do not wish admission to the hospital until the day of operation; a major reason being shorter hospital stay.[47] An updated systematic review on the effectiveness of 76 nurse-led preoperative assessment services for elective surgery found that the included 77 78 articles demonstrated a reduced length of stay; these studies had low methodological 79 quality, and therefore, the authors could not conclude that this service leads to reduced 80 length of hospital stay.[17] 381

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### 382 Strengths and limitations of the study

The review was performed in duplicate or independently by two researchers, and consensus 83 84 was reached through discussion. However, grey literature, such as government and institutional documents, was excluded and might be a limitation of this study. Since 85 organisation of healthcare systems varies among countries, the type of staff who performed 86 87 the preoperative assessment was not considered as an inclusion criterion. The European 88 Society of Anaesthesiology guidelines recommend that the anaesthesiologists must 89 complete the preoperative assessment while trained nurses or anaesthesia trainees should 90 perform the screening. [8] A preoperative evaluation performed by an internist was 91 associated with increased length of stay and increased postoperative mortality.[48] The 92 results of this systematic review may have been affected by the heterogeneity in the types 93 of staff performing the preoperative assessment. 94

We exclusively included studies with high internal validity. Therefore, several retrospective 95 96 studies were excluded. Nonetheless, as the remaining studies' risk of bias was fairly high, 97 and they were heterogeneous, a meta-analysis was not statistically appropriate. [26] The 98 included studies' designs could not rule out selection bias and confounders; thus, the 99 strength of the evidence should be assessed cautiously. Many studies did not adjust for 00 several confounders, which could be responsible for the observed effects. Several studies lacked descriptions of the methods used and the patients included, lowering transparency. 01 02 The results are relevant to health care services, focusing on the well-being and safety of the 23 patients. 04

#### 405 Implications for future research and practice

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406	This systematic review identified the ambiguity in the PAC interventions offered to the
407	intervention group. In many studies, it was evident that the methods used lacked clarity, and
408	high-quality research is needed in this field. The included studies did not demonstrate earlier
409	surgical room entry time[49,50] or reduction in the number of preoperative tests for
410	patients attending the PAC, similar to the results of the retrospective studies.[27]
411	Other implications may include the organisation structure of different PACs and their
412	functioning. Additionally, the tests that should be part of the assessment at the PACs should
413	be investigated. The use of technology, such as streaming services, facilitates different
414	patient groups and might become crucial for reducing human contact and spread of
415	infection in context of coronavirus disease 2019.
416	CONCLUSION
417	PAC use has reduced the length of stay and surgery cancellation rate at hospitals. However,
418	the effectiveness of PAC, the major review question, remains unclear, and requires further
419	research. There is a demand for high-quality studies capturing robust data describing the
420	quality of care and clinical outcomes for patients requiring anaesthesia. This requires
421	increased focus and funding for this specific area of health services research and could,
422	therefore, lead to implementation of PACs in health care services and improve patient safety
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428	The authors declare that they have no conflict of interest.
429	Data sharing statement

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27 28	440	EWK, MF, AO, and RCB: Quality assessment of the included articles				
29 30 31	441	EWK: Drafting of the manuscript				
32 33	442	MF, AO, RCB, and TOT: Contribution to and review of the final version of the manuscript				
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2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19	596	Figure legends
	597	
	598	Figure 1: Preferred Reporting Items for Systematic Reviews and Meta-Analysis (PRISMA).
	599	[23]
	600	Figure 2: The Joanna Briggs Institute Critical appraisal checklist for quasi-experimental
	601	studies was used for the risk of bias assessment. [29]
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Figure 2: The Joanna Briggs Institute Critical appraisal checklist for quasiexperimental studies was used for the risk of bias assessment. [30]

# Appendix 1: Search strategies

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### 1. Database Search Strategies

The search mode for CINAHL was Boolean/Phrase. For those searches that are done without search fields - it is automatically searched in the standard fields that CINAHL uses, including words from title, summary, and subject headings. Search 1 and Search 2 are with words in the title (TI in front of the keywords), however this have not been done with the other searches.

Embase and Medline have the same search mode because we are searching for words from title, summary, and subject headings.

#### 2. Initial searches, 11 September 2018

#### 2.1. Main Databases

Database: Embase, (Ovid)

Results: n=1287

Search:

1 ((pre-operativ\* or preoperativ\* or pre operativ\*) and (assessment\* or measurement\* or evaluat\*)).ti.

2 (clinic\* or unit\* or nurs\* or outpatient\* or ward\* or center\* or centre\*).ti.

3 1 and 2

4 ((preanaesthe\* or pre-anaesthe\* or pre anaesthe\* or pre-anesthe\* or preanesthe\* or pre anesthe\*) adj4 (assessment\* or measurement\* or evaluat\* or clinic\* or nurs\* or unit\* or outpatient\* or ward\* or center\* or centre\*)).ti,ab. (689)
5 (((pre-admiss\* or preadmiss\*) adj4 (assessment\* or measurement\* or evaluat\*

or clinic\* or unit\* or nurs\* or outpatient\* or ward\* or centre\* or center\*)) and (surg\* or anaesthe\* or anesthe\* or preoperativ\* or pre-operativ\* or preanaesthe\* or preanaesthe\* or pre anaesthe\* or pre-anesthe\* or preanesthe\* or preanesthe\*)).ti,ab.

6 ((anaesthe\* or anasthe\* or anesthe\*) adj4 outpatient\* adj4 clinic\*).ti,ab.

7 ((pre-admiss\* or preadmiss\*) adj4 (center\* or centre\*) adj4 (evaluat\* or assessment\* or measurement\*)).ti,ab.

- 8 or/3-7
- 9 limit 8 to yr="1996 -Current"
- 10 limit 9 to (conference abstracts or embase)

#### Database: Medline (Ovid)

Results: n=997

Search:

1 ((pre-operativ\* or preoperativ\* or pre operativ\*) and (assessment\* or measurement\* or evaluat\*)).ti.

2 clinic\* or unit\* or nurs\* or outpatient\* or ward\* or center\* or centre\*).ti.

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6	Clinic of nurs of unit of outpatient of ward of center of centre )).it,ab.
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14	7 ((pre-admiss <sup>*</sup> ) or preadmiss <sup>*</sup> ) adj4 (center <sup>*</sup> or centre <sup>*</sup> ) adj4 (evaluat <sup>*</sup> or
15	assessment* or measurement*)).ti,ab.
16	8 or/3-7
17	9 limit 8 to yr="1996 -Current"
18	10 limit 9 to (conference abstracts or embase)
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20	
22	Database: Medline (Ovid)
23	
24	Results: n=1105
25	Search:
26	1 ((pre-operativ* or preoperativ* or pre operativ*) and (assessment* or
27	measurement* or evaluat*)).ti.
20 20	2 (clinic* or unit* or nurs* or outpatient* or ward* or center* or centre*).ti.
30	3 1 and 2
31	4 ((preanaesthe* or pre-anaesthe* or pre anaesthe* or pre-anesthe* or
32	preanesthe* or pre anesthe*) adi4 (assessment* or measurement* or evaluat* or
33	clinic* or nurs* or unit* or outpatient* or ward* or center* or centre*)).ti.ab.
34	5 (((pre-admiss* or preadmiss*) adi4 (assessment* or measurement* or evaluat*
35	or clinic* or unit* or nurs* or outpatient* or ward* or centre* or center*)) and (surg*
30	or anaesthe* or anesthe* or preoperativ* or pre-operativ* or preanaesthe* or pre-
38	anaesthe* or pre anaesthe* or pre-anesthe* or preanesthe* or pre
39	anesthe*)) ti ab.
40	6 ((anaesthe* or anasthe* or anesthe*) adi4 outpatient* adi4 clinic*) ti ab.
41	7 ((pre-admiss* or preadmiss*) adi4 (center* or centre*) adi4 (evaluat* or
42	assessment* or measurement*)).ti.ab.
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55 54	measurement <sup>*</sup> or evaluat <sup>*</sup> ))
55	2 II (clinic" or unit" or nurs" or outpatient" or ward" or center or centre")
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57	4 (preanaestne <sup>*</sup> or pre-anaesthe <sup>*</sup> or "pre anaesthe <sup>*</sup> " or pre-anesthe <sup>*</sup> or
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5 ((pre-admiss\* or preadmiss\*) N3 (assessment\* or measurement\* or evaluat\* or clinic\* or unit\* or nurs\* or outpatient\* or ward\* or centre\* or center\*)) AND (surg\* or anaesthe\* or anesthe\* or preoperativ\* or "pre operativ\*" pre-operativ\* or preanaesthe\* or pre-anaesthe\* or pre-anesthe\* or preanesthe\* or "pre anesthe\*") 6 (anaesthe\* or anasthe\* or anesthe\*) N3 outpatient\* N3 clinic\* 23 (pre-admiss\* or preadmiss\*) N3 (center\* or centre\*) N3 (evaluat\* or assessment\* or measurement\*) 7 S3 OR S4 OR S5 OR S6 OR S7 487

8 S3 OR S4 OR S5 OR S6 OR S7 Limiters - Published Date: 19960101-; Exclude MEDLINE records

# 4. Search update, 4 February 2021

4.1. Main Databases

Database: Embase (Ovid)

Results: n=1572

Search:

1 ((pre-operativ\* or preoperativ\* or pre operativ\*) and (assessment\* or measurement\* or evaluat\*)).ti.

2 (clinic\* or unit\* or nurs\* or outpatient\* or ward\* or center\* or centre\*).ti.

3 1 and 2

4 ((preanaesthe\* or pre-anaesthe\* or pre anaesthe\* or pre-anesthe\* or preanesthe\* or pre anesthe\*) adj4 (assessment\* or measurement\* or evaluat\* or clinic\* or nurs\* or unit\* or outpatient\* or ward\* or center\* or centre\*)).ti,ab.

5 (((pre-admiss\* or preadmiss\*) adj4 (assessment\* or measurement\* or evaluat\* or clinic\* or unit\* or nurs\* or outpatient\* or ward\* or centre\* or center\*)) and (surg\* or anaesthe\* or anesthe\* or preoperativ\* or pre-operativ\* or preanaesthe\* or pre-anaesthe\* or pre-anaesthe\* or pre-anesthe\* or pre-anest

6 ((anaesthe\* or anasthe\* or anesthe\*) adj4 outpatient\* adj4 clinic\*).ti,ab.

7 ((pre-admiss\* or preadmiss\*) adj4 (center\* or centre\*) adj4 (evaluat\* or assessment\* or measurement\*)).ti,ab.

8 or/3-7

9 limit 8 to yr="1996 -Current"

10 limit 9 to (conference abstracts or embase)

Database: Medline (Ovid)

Results: n=1200

Search:

1 ((pre-operativ\* or preoperativ\* or pre operativ\*) and (assessment\* or measurement\* or evaluat\*)).ti.

2 (clinic\* or unit\* or nurs\* or outpatient\* or ward\* or center\* or centre\*).ti.

3 1 and 2

4 ((preanaesthe\* or pre-anaesthe\* or pre anaesthe\* or pre-anesthe\* or preanesthe\* or preanesthe\*) adj4 (assessment\* or measurement\* or evaluat\* or clinic\* or nurs\* or unit\* or outpatient\* or ward\* or center\* or centre\*)).ti,ab.

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# Reporting checklist for systematic review and meta-analysis.

Based on the PRISMA guidelines.

# Instructions to authors

Complete this checklist by entering the page numbers from your manuscript where readers will find each of the items listed below.

Your article may not currently address all the items on the checklist. Please modify your text to include the missing information. If you are certain that an item does not apply, please write "n/a" and provide a short explanation.

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In your methods section, say that you used the PRISMAreporting guidelines, and cite them as:

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		Page
	Reporting Item	Number
Title	4	
<u>#1</u>	Identify the report as a systematic review, meta-analysis, or both.	1
Abstract		
Structured <u>#2</u> summary	Provide a structured summary including, as applicable: background; objectives; data sources; study eligibility criteria, participants, and interventions; study appraisal and synthesis methods; results; limitations; conclusions and implications of key findings; systematic review registration number	2
Introduction		
Rationale <u>#3</u>	Describe the rationale for the review in the context of what is already known.	3-5
Foi	r peer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml	

1 2 3 4 5	Objectives	<u>#4</u>	Provide an explicit statement of questions being addressed with reference to participants, interventions, comparisons, outcomes, and study design (PICOS).	5	
6 7	Methods	lethods			
8 9 10 11 12	Protocol and registration	<u>#5</u>	Indicate if a review protocol exists, if and where it can be accessed (e.g., Web address) and, if available, provide registration information including the registration number.	5	
14 15 16 17 18 19	Eligibility criteria	<u>#6</u>	Specify study characteristics (e.g., PICOS, length of follow- up) and report characteristics (e.g., years considered, language, publication status) used as criteria for eligibility, giving rational	6	
20 21 22 23 24 25 26	Information sources	<u>#7</u>	Describe all information sources in the search (e.g., databases with dates of coverage, contact with study authors to identify additional studies) and date last searched.	5,6	
27 28 29 30 31	Search	<u>#8</u>	Present full electronic search strategy for at least one database, including any limits used, such that it could be repeated.	5,6 + APPENDIX 1	
32 33 34 35 36 37 38	Study selection	<u>#9</u>	State the process for selecting studies (i.e., for screening, for determining eligibility, for inclusion in the systematic review, and, if applicable, for inclusion in the meta-analysis).	6	
39 40 41 42 43 44 45	Data collection process	<u>#10</u>	Describe the method of data extraction from reports (e.g., piloted forms, independently by two reviewers) and any processes for obtaining and confirming data from investigators.	6-7	
46 47 48 49 50	Data items	<u>#11</u>	List and define all variables for which data were sought (e.g., PICOS, funding sources), and any assumptions and simplifications made.	5-7	
51 52 53 54 55 56 57 58 59 60	Risk of bias in individual studies	<u>#12</u> For p	Describe methods used for assessing risk of bias in individual studies (including specification of whether this was done at the study or outcome level, or both), and how this information is to be used in any data synthesis.	6-7	

1 2 3	Summary measures	<u>#13</u>	State the principal summary measures (e.g., risk ratio, difference in means).	N/A	
4 5 6 7 8	Planned methods of analyis	<u>#14</u>	Describe the methods of handling data and combining results of studies, if done, including measures of consistency (e.g., I2) for each meta-analysis.	7	
10 11 12 13 14	Risk of bias across studies	<u>#15</u>	Specify any assessment of risk of bias that may affect the cumulative evidence (e.g., publication bias, selective reporting within studies).	12	
15 16 17 18 19 20	Additional analyses	<u>#16</u>	Describe methods of additional analyses (e.g., sensitivity or subgroup analyses, meta-regression), if done, indicating which were pre-specified.	N/A	
20	Results				
22 23 24 25 26 27	Study selection	#17 Give numbers of studies screened, assessed for eligibility, and included in the review, with reasons for exclusions at each stage, ideally with a <u>flow diagram</u> .			
28 29 30 31 32	Study characteristics	<u>#18</u>	18 For each study, present characteristics for which data were extracted (e.g., study size, PICOS, follow-up period) and provide the citation.		
33 34 35 36	Risk of bias within studies	<u>#19</u>	Present data on risk of bias of each study and, if available, any outcome-level assessment (see Item 12).	12 + Figure	
<ol> <li>37</li> <li>38</li> <li>39</li> <li>40</li> <li>41</li> <li>42</li> <li>43</li> </ol>	Results of individual studies#20For all outcomes considered (benefits and harms), present, for each study: (a) simple summary data for each intervention group and (b) effect estimates and confidence intervals, ideally with a forest plot.		12-16		
44 45 46 47 48	Synthesis of#21Present the main results of the review. If meta-analyses are done, include for each, confidence intervals and measures of consistency.		11-16		
49 50	Risk of bias	<u>#22</u>	Present results of any assessment of risk of bias across	N/A	
51 52	across studies		studies (see Item 15).		
53 54	Additional	<u>#23</u>	Give results of additional analyses, if done (e.g., sensitivity	N/A	
55 56	analysis		or subgroup analyses, meta-regression [see Item 16]).		
57 58	Discussion				
5960For peer review only - http://bmjopen.bmj.com/site/about/gui			eer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml		

1 2 3 4 5 6	Summary of Evidence	<u>#24</u>	Summarize the main findings, including the strength of evidence for each main outcome; consider their relevance to key groups (e.g., health care providers, users, and policy makers	16-19
7 8 9 10 11	Limitations	<u>#25</u>	Discuss limitations at study and outcome level (e.g., risk of bias), and at review level (e.g., incomplete retrieval of identified research, reporting bias).	19
12 13 14 15	Conclusions	<u>#26</u>	Provide a general interpretation of the results in the context of other evidence, and implications for future research.	20
17 18	Funding			
19 20 21 22 23	Funding	<u>#27</u>	Describe sources of funding or other support (e.g., supply of data) for the systematic review; role of funders for the systematic review.	21
24 25	None The PRISMA	A check	klist is distributed under the terms of the Creative Commons A	ttribution
26 27	License CC-BY. Th	nis che	cklist can be completed online using <u>https://www.goodreports.</u>	<u>org/</u> , a tool
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# **BMJ Open**

# Effectiveness of pre-anaesthetic assessment clinic: A systematic review of randomised and non-randomised prospective controlled studies

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<b>Primary Subject Heading</b> :	Anaesthesia
Secondary Subject Heading:	Surgery, Nursing, Health services research
Keywords:	Adult anaesthesia < ANAESTHETICS, Health & safety < HEALTH SERVICES ADMINISTRATION & MANAGEMENT, Quality in health care < HEALTH SERVICES ADMINISTRATION & MANAGEMENT, Adult surgery < SURGERY

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2 3 4	1	Effectiveness of pre-anaesthetic assessment clinic: A systematic review of randomised and
5 6 7	2	non-randomised prospective controlled studies
7 8 9	3	
10 11	4	Eirunn Wallevik Kristoffersen <sup>1,2</sup> *, Anne Opsal <sup>1</sup> , Tor Oddbjørn Tveit <sup>1,2,3</sup> , Rigmor C Berg <sup>4,5</sup> ,
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49 50 51	20	
52 53	21	Word count: 4156
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3 4	24	ABSTRACT
5 6 7	25	Objectives: The aim of this systematic review was to examine the effectiveness of pre-
	26	anaesthesia assessment clinics (PACs) in improving the quality and safety of perioperative
8 9	27	patient care.
10 11	28	Design: Systematic review.
12 13	29	Data sources: The electronic databases CINAHL Plus with Full Text (EBSCOhost), Medline,
14 15	30	and Embase (OvidSP) were systematically searched on 11 September 2018
16	31	and updated on 3 February 2020 and 4 February 2021.
17	32	Eligibility criteria: The inclusion criteria for this study were studies published in English or
19 20	33	Scandinavian language and scientific original research that included randomised or non-
21 22	34	randomised prospective controlled studies. Additionally, studies that reported the outcomes
23 24	35	from a PAC consultation with the patient present were included.
25	36	Data extraction and synthesis: Titles, abstracts, and full texts were screened by a team of
26 27	37	three authors. Risk of bias was assessed using the Joanna Briggs Institute critical appraisal
28 29	38	checklist for quasi-experimental studies. Data extraction was performed by one author and
30 31 32 33	39	checked by four other authors. Results were synthesised narratively owing to the
	40	heterogeneity of the included studies.
34 35	41	Results: Seven prospective controlled studies on the effectiveness of PACs, were included.
36 37	42	Three studies reported a significant reduction in the length of hospital stay and two studies
38	43	reported a significant reduction in cancellation of surgery for medical reasons when patients
39 40	44	were seen in the PAC. In addition, the included studies presented mixed results regarding
41 42	45	anxiety in patients. Most studies had a high risk of bias.
43 44	46	Conclusion: This systematic review demonstrated a reduction in the length of hospital stay
45 46	47	and cancellation of surgery when the patients had been assessed in the PAC. There is a need
47	48	for high-quality prospective studies to gain a deeper understanding of the effectiveness of
48 49	49	PACs.
50 51	50	PROSPERO registration number: CRD42019137724
52 53	51	
54 55	52	Keywords: pre-anaesthetic assessment clinic, preoperative care, quality, safety, systematic
56 57	53	review
57	54	
59 60	55	

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2 3	56	
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8 9	60	Strengths and limitations of this study
10 11	61	An extensive database search was conducted with no limitations on outcomes and
12	62	the type of pre-anaesthetic assessment clinic.
13 14 15	63	Only randomised or non-randomised prospective controlled studies were included.
15 16	64	• The Joanna Briggs Institute Critical appraisal checklist for quasi-experimental studies
17 18	65	was used.
19 20	66	• The included studies were heterogeneous and had a high risk of bias, which is a
21 22	67	major limitation of this review.
23 24	68	
2 <del>4</del> 25 26	69	INTRODUCTION
20 27 28	70	Anaesthesia is crucial in surgery. However, it may activate physiological changes that
29 30 31	71	increase morbidity and mortality,[1] depending on the patients' preoperative health status
32 33	72	and age.[2] Hospitals treat patients with complex, comorbid healthcare problems,
34 35 36	73	undergoing progressively extensive surgeries and interventions.[3,4] To ensure the quality
37 38	74	and safety of anaesthesia and surgery, precise knowledge of the clinical characteristics of
39 40	75	patients is critical to the perioperative management.[2] Over the past 50 years,
41 42 43	76	perioperative mortality, including anaesthesia-related mortality, has declined, which is
44 45	77	significant in developed countries,[1,5] mainly because of new anaesthetics, improved
46 47 48	78	monitoring equipment and training, availability of recovery rooms, and improved airway
49 50	79	management.[4] However, a previous review found higher rates of morbidity and mortality
51 52 53	80	in non-operating room anaesthesia, which was attributed to limited preoperative
54 55	81	evaluation.[6] A retrospective study found significant associations between perioperative
56 57	82	mortality and age < 1 year or > 65 years, American Society of Anesthesiologists Physical
59 60	83	Status Classification System (ASA), emergency case status, and operative start time after

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84 6:00 PM.[7] This might indicate that risk factors are both patient- and surgery-related and 85 may be linked to organisational structures.[8]

86 Currently, an increasing number of pre-anaesthesia assessment clinics (PACs) support 87 hospitals internationally in handling the rising number of patients and complexity of surgical 88 procedures.[9] The design of PACs differs critically based on location, organisational 89 structure, timing, and patient groups. They primarily function as a service unit for surgeons, 90 patients, and the anaesthetic team.[10] The PAC consultation, by the anaesthesiologist, 91 anaesthesia nurse, or both, is a globally recognised evaluation method and optimises the 92 patients' medical condition prior to surgery and anaesthesia.[2] Thus, it is essential for 93 secure anaesthetic practice since it detects anaesthesia-related risk factors and high-risk 94 patients, improves patient outcomes, prepares the patient physically and psychologically for 95 anaesthesia, and ensures the patient's most favourable condition for surgery and 96 anaesthesia.[11-13] This is primarily performed by interviewing and examining the patient; 97 reviewing previous medical, surgical and anaesthesia issues; evaluating current medication; 98 and obtaining and reviewing preoperative tests.[10] PACs also allow increased 99 communication between healthcare providers and coordination with postoperative .00 care.[14,15] Because of well-prepared patients and staff, several researchers have indicated .01 that with PAC, the number of surgical cancellations, length of hospital stay, laboratory tests, .02 and mortality rate have reduced. [7,16,17] Others assert that patients feel less anxious .03 regarding the subsequent anaesthetic and surgical processes and are highly satisfied with .04 the service with PAC consultations.[16,18,19]

.05 As Turunen et al. stated, research on PACs regarding costs, financial savings, the impact on .06 patient safety and quality of care, accuracy of the number of operative patients, and effect

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107 on preoperative nursing levels, is scarce.[20] Survey results indicate that anaesthesiologists
108 perceive day of surgery delays due to missing information as common, even with PAC
109 consultations.[21] This systematic review aimed to examine the effectiveness of PACs in
110 improving the quality and safety of perioperative patient care. Further, we aimed to
111 determine the gaps in existing knowledge for future research.
112 METHODS

Our systematic review followed the guidelines in the Cochrane Handbook for Systematic
 Reviews of Interventions[22] and was reported in accordance with the Preferred Reporting
 Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines.[23] The protocol was
 registered in PROSPERO: CRD42019137724.[24]

9 117 The two review questions were:

118 1. Is PAC effective in improving patient satisfaction and safety, while reducing anxiety?

119 2. Is PAC effective in reducing cancellation rate and cost of surgery, and improving the

120 efficiency of perioperative patient care?

#### 121 Search strategies

122 We performed a scoping search in different databases to identify the key terms.[25,26] The

123 final search was planned and conducted in close collaboration with the university librarian.

124 On 11 September 2018 we searched CINAHL Plus with Full Text (EBSCOhost), Medline, and

125 Embase (OvidSP) databases, which were updated on 3 February 2020 and 4 February 2021.

126 Considering the lack of subject headings (e.g., MeSH) for PAC, we combined text words, such

- 127 as 'pre-anaesthesia', 'nurse', 'surgery', 'anaesthesia', 'preoperative', 'assessment',
- 128 'measurement', 'evaluate', 'preadmission', 'centre', 'clinic', 'ward', 'unit', and 'outpatient'.
- 129 The searches are detailed in Appendix 1. The search mode in CINAHL was Boolean/Phase,

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3 4	130	which supports Boolean searching or exact phrase searching. For comprehensiveness, we
5 6 7	131	used both the truncation and proximity operators. We limited the search to 1996, the year
7 8 9	132	one of the first known articles in this area was published.[27] Complementary methods to
10 11	133	identify studies included following up on citations via Scopus, scanning the reference lists of
12 13 14	134	relevant papers and included articles, and checking for relevant studies in clinical trials.[25]
15 16	135	Eligibility criteria
17 18 19	136	The main inclusion criterion was that the study, using empirical quantitative methods,
20 21	137	addressed the effectiveness of PACs. Specific eligibility criteria were: (a) published in English
22 23 24	138	or Scandinavian language, (b) scientific publication of original research, (c) reported the
25 26	139	outcomes of PAC, (d) PAC consultation with the patient present, (e) randomised or non-
27 28 29	140	randomised prospective controlled studies, and (f) newly established PAC. The following
30 31	141	were excluded: (a) editorials, discussions, and conference abstracts, (b) reviews, (c)
32 33 34	142	instrument testing, (d) studies on children, and (e) retrospective studies.
35 36	143	Study selection
37 38 30	144	All references identified in the search were transferred to EndNoteX9, where the duplicates
39 40 41	145	were removed. Subsequently, all unique references were transferred to the Covidence
42 43	146	screening tool.[28] Study eligibility was ascertained independently, first at the title and
44 45 46	147	abstract level, and subsequently at the full text level. Three of the authors screened all the
47 48	148	articles (EWK, AO, MF). Inclusion was determined by consensus, and disagreements were
49 50 51	149	resolved by consulting two other authors (RCB and TOT).
52 53	150	Quality assessment
54 55 56	151	Risk of bias in studies were assessed using design-specific checklists. Given the
57 58	152	methodological similarity of the studies, only the Joanna Briggs Institute Critical appraisal
59 60	153	checklist for quasi-experimental studies was used.[29] Author EWK performed the risk of

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154 bias assessment, and RCB confirmed its accuracy. Disagreements were resolved through .55 discussion with MF and AO. Each of the nine checklist questions was answered no, yes, unclear, or not applicable. .56

.57 Data extraction and analysis

Author EWK extracted data from each study onto a pre-designed Excel spreadsheet. All the .58 .59 authors confirmed the accuracy, consistency, and completeness of the extracted data that .60 included publication details; study design; setting; and characteristics of the patients, .61 interventions, comparisons, and outcome (PICO). We requested information on the missing data; however, received no response from the authors. If the PICO elements had been .62 .63 sufficiently similar and statistical data were available, we had intended to conduct a meta-.64 analysis. However, the extracted data revealed substantial heterogeneity. Therefore, we .65 performed a narrative synthesis. We described the findings in text, stratified by outcome, with descriptions of the effects of interventions for each study, classification of the effect .66 .67 direction, and we looked across contributing studies to develop a summary of findings for .68 each outcome.[22] .69 **Patient and Public Involvement** .70 Patients and/or the public were not involved in the design, conduct, reporting, or .71 dissemination plans of this research. However, the project was initiated by health .72 professionals.

.74 RESULTS

> .75 Figure 1. provides the details of the study selection process. A total of 2,981 records were .76 identified in the final search (2021). After removing duplicates, we screened 2,058 records

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The seven studies are listed in Table 1. All seven studies were in English and were published

report data collection information).[30] Based on the inclusion criteria, all were prospective

between 2000 and 2017, with data collected between 1997 and 2015 (one study did not

controlled studies; however, no RCTs were found. There was one controlled before-after

implementation, without baseline assessments. There were three two-group non-parallel

after-only studies, [32-34] and three two-group parallel after-only studies; [30] one had a

matched control group[35] and one had three follow-up assessments of one arm.[36] One

study had only cancellation rate as prospective data.[32] The studies included 77,411

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study.[31] The remaining six studies had control groups; assessments followed PAC

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patients.

based on the title and abstract; 179 records passed the full-text screening. After applying theinclusion criteria, seven studies were selected for the final analysis.

**Overall characteristics of the studies** 

#### **Table 1: Description of the included studies**

Author, Year, Country	Study design	Sampling time	Population	Intervention	Comparison	Outcomes
Farasatkish,2009[33] Iran	Two-group after study	May 2007 through August 2007	N=1716, open- heart surgery, ASA class III-IV	Pre-anaesthesia consultation clinic (3-10 days before surgery)	Usual care (within 24 h of surgery)	Cancellations
Kamal,2011[34] England	Two-group	April 2005 through April 2009	N=1445, complex elective orthopaedic surgery, ASA class III-IV	Preoperative anaesthetic assessment clinic (timing not stated)	Usual care (day of surgery)	Admissions, length of stay, mortality, cost
Kamau,2017[31] Kenya	СВА	August 2000, April 2001, November 2001	N=51, elective non-cardiac surgery, ASA class not stated	Pre-anaesthesia clinic consultation (≥48 h before surgery)	Usual care (day before surgery)	Anxiety (STAI score)
Klopfenstein,2000[30] Switzerland	Two-group after study (parallel)	No data	N=40, elective endoscopic urological surgery, ASA class I-III	Pre-anaesthetic consultation (1-2 weeks before surgery)	Usual care (the evening before surgery)	Anxiety (MAACL, VAS)
Lee,2012[35] China	Two-group after study (parallel)	March 2007 through November 2009	N=352, elective surgery, ASA class I-IV	Anaesthesia consultation clinic (≤3 months before surgery)	Usual care (the evening before surgery)	Quality of recovery score, cost, cancellations, length of stay, satisfaction, anxiety (VAS), willingness to pay (WTP)
Mendes,2005[36] Brazil	Two-group after study (parallel)	April 2007 through June 2007	N=52254, surgery, ASA class not stated	Preoperative outpatient evaluation clinic (timing not stated)	Usual care (timing not stated)	Cancellations, length of stay

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2 3 4 5 6 7	101	van Klei,2002[32] The Netherlands	Two-group after study	November 2012	N=21553, elective surgery, ASA class not stated	Preoperative outpatient evaluation clinic (average 3 weeks before surgery)	Usual care (day before surgery)	Cancellations
8 9	191							
10	192	ASA: American Societ	y of Anesthesiol	ogy; CBA: controlle	ed before-after; M	AACL: Multiple Affect Adj	ective Check List; S	TAI: State-Trait Anxiety
12 13 14 15 16 17	193	Inventory; VAS: visua	l analogue scale;	WTP: willingness	to pay			
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194	Of the 77,411 patients in the studies, 9,626 and 15,531 patients were in the intervention and
195	control groups, respectively. One study did not specify the number of patients in the
196	intervention and control groups, but only the total number of surgeries performed.[36] Five
197	studies reported data for sex, showing that 51% of the patients were women and 49% were
198	men (12,129 vs. 11,583).[30-33,35] There were more women than men in both the
199	intervention (4,345 vs. 4,134) and control groups (7,784 vs. 7,449). Five studies reported
200	data for age showing that all patients were over 20 years of age[30-33,35] and four studies
201	had grouped within the American Society of Anesthesiology (ASA) category.[30,33-35]
202	
203	The patients were scheduled to undergo a variety of surgeries, including
204	orthopaedic,[31,32,34,35] urology,[30-32,35] general,[31,32,35] heart,[33]
205	gynaecology/obstetrics,[31,32,35] vascular surgery,[32] ophthalmology,[32]
206	maxillofacial/dental surgery,[31,32] and neurological surgery,[32] while one did not specify
207	the type.[36] In five studies, the type of anaesthesia was not specified,[31-34,36] and two
208	studies reported patients for general and/or regional supplement.[30,35]
209	
210	The patients included had previous anaesthetic experience in one study,[30] previous and no
211	previous anaesthetic experience in another,[31] and five studies did not report this data.[32-
212	36] Limited background characteristics of the patients were reported in two studies.[34,36]
213	One stated that the patients included had ASA 3 or 4 and a body mass index >40; however,
214	no ASA number, sex, or age was reported.[34] Mendes et al. did not report any background
215	characteristics of the included patients.[36]
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217	Considering the intervention, the PACs in all studies comprised an outpatient service
218	whereby patients were examined for medical conditions important for anaesthesia and
219	informed regarding expectations on the day of surgery. Nevertheless, the terminology used
220	for PACs varied, as they served different surgical specialities and conducted pre-anaesthesia
221	consultation from $\geq$ 48 h to $\leq$ 3 months before the surgery. The settings included a university
222	hospital (n=3),[31,35,36] teaching hospital (n=1),[32] medical centre (n=1),[33] and general
223	hospital (n=1);[34] one study did not specify the context.[30] The staff conducting the pre-
224	anaesthetic consultation also varied: in five studies, it was the anaesthesiologists,[30,32,34-
225	36] in the other studies, it was (also) the orthopaedic senior house officer,[34] consultant or
226	resident,[31] or physician.[33] In three studies, nurses were part of the team.[32,34,35] The
227	comparison in all studies was usual care, which generally involved a preoperative
228	anaesthetic evaluation of the admitted patients the day before the surgery.
229	Description of risk of bias in the studies
230	Figure 2. shows the results of the risk of bias assessment. In all seven included studies, the
231	cause and effect were clear. Most of the studies measured outcomes similarly and used
232	appropriate statistical analyses. Several studies had limitations of follow-up and similarity in
233	care and participants. None of the patients had multiple pre-and post-measurements.
234	Outcomes of the included studies
235	The outcomes of the included studies are each described separately below.
236	Satisfaction
237	One study reported satisfaction as an outcome.[35] The summarised patient satisfaction
238	with the anaesthetic consultation score out of 100 showed that the patients in the PAC
239	group were more satisfied (mean difference, 2.10%; 95% confidence interval [CI], 0.51–
240	3.70%; <i>p</i> =0.01).[35] There was no statistically significant difference between the two groups

in mean patient satisfaction with perioperative anaesthesia care score after surgery (mean difference 0.01%, p=0.94).[35] The quality of recovery (QoR) measure referred to the patients' quality of recovery score.[37] The mean QoR score (range, 0–18) following anaesthesia on the first day after surgery was similar between the intervention (13.17±2.73) and control (13.31±2.65) groups (*p*=0.67).[35] Anxiety Three studies reported anxiety. [30,31,35] Two studies reported the visual analogue scale (VAS), one rated from zero (no anxiety) to ten (very high anxiety),[30] another used a 100-mm horizontal line with 'not anxious at all' to 'extremely anxious'.[35] In one study, the median VAS anxiety score was 3 (0–5) in the intervention group and 5 (2–8) in the control group (p=0.0038).[30] In another study, there were no significant differences between the control and intervention groups for levels of anxiety (VAS), surgery (26 vs. 25, respectively, p=0.12), and anaesthesia (20 vs. 19, respectively, p=0.60).[35] The median Multiple Affect Adjective Check List (MAACL) score, with possible range of scores from 0 to 21 (higher scores indicating greater levels of anxiety), was 3 (0–9) in the intervention group and 6.5 (2–12) in the control group (p=0.0053).[30] The differences in the State-Trait Anxiety Index (STAI) score, which comprised 40 questions rated on a four-point Likert scale, was 1.51 (95% CI: 1.02–2.02%, p=0.0051).[31] The results on anxiety in these two studies were significant. However, Kamau et al. found no differences on examining anxiety and the influences of sex, duration of hospital stay, and prior anaesthesia experience.[31] Mortality One study reported the mortality rates.[34] Patients attending the High Dependency Unit (HDU), Intensive Care Unit (ICU), and Post-anaesthesia Care Unit (PACU) following complex 

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264	orthopaedic surgery had a significant reduction in mortality rate after being assessed at the
265	PAC, from 18 (6.1%) of 298 patients to 14 (1.2%) of 1147 patients ( <i>p</i> =0.001).[34]
266	Cancellation rate
267	Four studies reported reduced cancellation rates following the establishment of a
268	PAC.[32,33,35,36] One of the included studies had 316 (2.0%) cancellations for medical
269	reasons before the introduction of PAC, and 79 (0.9%) after, with a difference of 1.02% (95%
270	CI, 0.31–1.31%). After adjustment, the odds ratio was 0.7 (95% CI, 0.5–0.9%).[32] The overall
271	cancellation of surgery reduced from 1027 (6.3%) to 393 (4.6%) following PAC introduction,
272	with a difference of 0.9% (95% Cl, 0.3–1.0%).[32] Mendes <i>et al</i> .[36] found a decrease in
273	overall cancellations from year 1 (39.3%) to year 4 (15.9%), $p \le 0.05$ . In the first year of their
274	study, there were 469 cancellations per 10,639 surgeries performed. The following year, a
275	considerable increase above the baseline in the intervention group was observed, followed
276	by a progressive decrease in the last year with 391 cancellations per 10,397 surgeries
277	performed.[36] Farasatkish et al. reported that of the 1,716 patients studied, a mean of 15.1
278	% cancelled in the two groups. The cancellation rates in the control and intervention groups
279	were 16.8% (146 [number of cancellations]/866 [number of surgeries]) and 13.29%
280	(113/850) (p=0.046), respectively. The most common reason for cancellation was
281	incomplete medical work-up; 51/146 (35%) in the control group and 32/113 (28%) in the
282	intervention group (p=0.03).[33] Lee et al. found similar rates for surgery being cancelled on
283	the scheduled date for the intervention group compared with the control group (2.3% vs.
284	3.4%, <i>p</i> =0.75).[35]
285	
286	Costs and willingness to pay

3 4	287	Two studies reported the costs.[34,35] One study reported a total saving of £ 486.62 per
5 6 7	288	patient after establishing a PAC.[34] Another reported a significantly lower preoperative cost
7 8 9	289	per patient in the intervention group compared with that of the control group (mean
10 11	290	difference, \$ 463; 95% Cl, -\$648 to -\$278 per patient, <i>p</i> <0.01).[35] However, the mean
12 13 14	291	difference in the total perioperative treatment cost was not significant, even after adjusting
14 15 16	292	for cancellation on the day of surgery costs.[35] Compared to the control group, the
17 18	293	willingness to pay (WTP) among the intervention group patients was significantly more than
19 20 21	294	the median WTP (US \$13) for a clinic consultation at the PAC.[35]
22 23	295	Length of stay
24 25 26	296	The length of stay was reported in three studies.[34-36] Mendes et al.[36] found a significant
27 28	297	decrease in mean hospital stay of patients from 6.2 to 5.0 days (p $\leq$ 0.001) during the 4 years
29 30 31	298	of this study. Kamal <i>et al.</i> [34] found a significant reduction in the length of stay in the HDU
32 33	299	from 2.1 days to 1.6 days ( $p$ =0.01), and in the ICU from 2.3 days to 1.9 days ( $p$ =0.01). In the
34 35	300	last study, no significant changes were found in the median duration of postoperative stay
36 37 38	301	between the intervention and control groups.[35]
39 40	302	Organisation planning and efficiency
41 42 43	303	Organisation planning and efficiency have been reported in two studies.[34,36] One study
44 45	304	found statistically significant changes in the reduction of unplanned admissions to the PACU
46 47 48	305	(65/298 [22%], 111/1147 [10%], <i>p</i> =0.001), ICU (4/298 [1.3%], 4/1147 [0.4%], <i>p</i> =0.01), and
49 50	306	HDU (4/298 [1.34%], 20/1147 [1.7%], <i>p</i> =0.01) after implementing a PAC.[34] The planned
51 52	307	admissions in the ICU (4/298 [1.3%], 18/1147 [1.6%], <i>p</i> =0.01), and HDU (14/298 [4.7%],
53 54 55	308	85/1147 [7.4%], <i>p</i> =0.1) increased after implementing a PAC.[34] The number of PAC
56 57	309	evaluations increased from 14,704 (year 1) to 413,990 (year 4) ( $p \le 0.001$ ).[36] The number
58 59 60		

)4	found statistically significant changes in the reduction of unplanned admissions to the PA	CU
)5	(65/298 [22%], 111/1147 [10%], $p$ =0.001), ICU (4/298 [1.3%], 4/1147 [0.4%], $p$ =0.01), and	
)6	HDU (4/298 [1.34%], 20/1147 [1.7%], <i>p</i> =0.01) after implementing a PAC.[34] The planned	
)7	admissions in the ICU (4/298 [1.3%], 18/1147 [1.6%], <i>p</i> =0.01), and HDU (14/298 [4.7%],	
)8	85/1147 [7.4%], <i>p</i> =0.1) increased after implementing a PAC.[34] The number of PAC	
)9	evaluations increased from 14,704 (year 1) to 413,990 (year 4) ( $p \le 0.001$ ).[36] The number	r
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3 4	310	of outpatient procedures increased from 2,170 (year 1) to 1,943 (year 4) ( $p \le 0.001$ ) and the
5 6 7	311	inpatient procedures decreased from 9,556 (year 1) to 8,449 (year 4) ( $p \le 0.001$ ).[36]
<ul> <li>7</li> <li>8</li> <li>9</li> <li>10</li> <li>11</li> <li>12</li> <li>13</li> <li>14</li> <li>15</li> <li>16</li> <li>17</li> <li>18</li> <li>19</li> <li>20</li> </ul>	312 313	DISCUSSION
	314	This systematic review summarises the effectiveness of PACs in improving quality and safety
	315	of pre-anaesthetic patient care in general hospitals and determines the gaps in existing
	316	knowledge for future research. Herein, we present the main results of the review and infer
	317	the implications for research and practice.
21 22	318	
23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39	319	Seven studies met the inclusion criteria, and the main findings were reduction in the length
	320	of stay and surgery cancellation rate in hospitals. However, the studies were of low quality,
	321	making it difficult to draw any conclusion. The evidence from our systematic review is
	322	insufficient to conclude whether patients have reduced anxiety because of PAC. This is
	323	because the included studies used different instruments for measuring anxiety levels, and
	324	the results could not be pooled.[38]
	325	
40 41 42	326	A major purpose of establishing a PAC in a hospital is to better prepare the patients for the
43 44	327	anticipated surgery. Healthcare professionals and policymakers are exploring strategies to
45 46 47	328	reduce unnecessary investigations without compromising quality of care and patient
48 49	329	safety.[39] Transition of evidence-based interventions to the hospital systems can provide
50 51 52	330	substantial benefits to patient care.[40] According to the ASA and the National Institute of
52 53 54 55 56 57 58 59 60	331	Health and Care Excellence, routine preoperative laboratory tests are not recommended for
	332	relatively healthy patients. Instead, they encourage patient and surgery- specific
	333	investigations.[15,40] This recommendation is not always implemented in hospital protocols

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2 3	334	or practice. An observational study showed that routine preoperative testing to predict		
4 5 6	335	abnormalities found at least one abnormal test result in most of the relatively healthy		
7 8 9	336	patients. Only 0.67% of the abnormalities had a significant impact on changing the		
9 10 11 12 13 14	337	perioperative management.[41] Blitz et al. argued that PACs should focus on early patient		
	338	engagement strategies, interdisciplinary team communication, detailed perioperative care		
15 16	339	plans, and patient documentation using electronic health record, which should be open for		
17 18 19 20 21 22	340	review by the perioperative team.[14] Furthermore, a previous study mentioned that the		
	341	risk factors are not only patient-related but also organisation-related,[7] and that some		
22 23 24	342	hospitals have perioperative care teams that are better at identifying and relieving		
25 26 27 28 29 30 31 32 33 34 35 36	343	perioperative complications.[42,43] Thus, the value of PACs lies in their ability to improve		
	344	the quality of the perioperative process by designing a more robust system for preoperative		
	345	assessment and preparation.[14] A narrative review found higher rates of morbidity and		
	346	mortality in non-operating room anaesthesia, and one of the main reasons was limited		
	347	preoperative evaluation.[6] In this systematic review, the assessment of PAC was		
37 38 39	348	significantly associated with reduced mortality following complex orthopaedic surgery in		
40 41	349	only one study.[34] Retrospective studies have reported similar results, but with different		
42 43 44	350	surgeries.[14,44]		
45 46	351			
47 48 49	352	Cancellation on the day of surgery has undesirable effects on both the patients and the		
49 50 51 52 53 54 55 56	353	hospital system.[13] Late patient-related cancellations can totally or partially be		
	354	prevented, [45] if addressed during preoperative evaluations. [16] This has been confirmed by		
	355	only three studies in this systematic review that found a reduction in surgery cancellation		
57 58	356	after implementing a PAC.[32,33,36] However, Lee <i>et al.</i> found no significant changes		
60	357	between the intervention and control groups.[35] Mendes et al. found that the number of		

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358	cancellations for medical reasons after PAC implementation decreased in the first year of
359	implementation. In the second and third years, they were high before the number dropped
360	to below baseline.[36] These conflicting findings indicate that hospitals operate in specific
361	contexts, with unique populations, processes, and microsystems, encountering unique
362	obstacles, making implementation difficult. Patient-focused interventions should consider
363	barriers, facilitators, and interrelationships between systems, staff, and interventions to
364	increase the likelihood of sustainable success.[46] Additionally, Kamal et al. indicated that
365	PACs lead to more planned admissions to the ICU, HDU, and PACU, which is more
366	predictable for patients, staff, and administration.[34]
367	
368	Another finding of this review was a significant reduction in the length of hospital stay
369	following patients' examination in a PAC; however, a small number of studies with low
370	quality were considered. Nevertheless, similar results were found in another systematic
371	review claiming that perioperative systems support the hospitals by addressing the expected
372	growth in the number and complexity of surgical procedures.[16] When patients are
373	examined in the PAC and well-prepared with information, consultations, and tests, they
374	need not be hospitalised until the day of surgery. A survey on operated patients showed that
375	given a choice, 75% do not wish admission to the hospital until the day of operation; a major
376	reason being shorter hospital stay.[47] An updated systematic review on the effectiveness of
377	nurse-led preoperative assessment services for elective surgery found that the included
378	articles demonstrated a reduced length of stay; these studies had low methodological
379	quality, and therefore, the authors could not conclude that this service leads to reduced
380	length of hospital stay.[17]
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#### 382 Strengths and limitations of the study

The review was performed in duplicate or independently by two researchers, and consensus 83 84 was reached through discussion. However, grey literature, such as government and institutional documents, was excluded and might be a limitation of this study. Since 85 organisation of healthcare systems varies among countries, the type of staff who performed 86 87 the preoperative assessment was not considered as an inclusion criterion. The European 88 Society of Anaesthesiology guidelines recommend that the anaesthesiologists must 89 complete the preoperative assessment while trained nurses or anaesthesia trainees should 90 perform the screening. [8] A preoperative evaluation performed by an internist was 91 associated with increased length of stay and increased postoperative mortality.[48] The 92 results of this systematic review may have been affected by the heterogeneity in the types 93 of staff performing the preoperative assessment. 94

We exclusively included studies with high internal validity. Therefore, several retrospective 95 96 studies were excluded. Nonetheless, as the remaining studies' risk of bias was fairly high, 97 and they were heterogeneous, a meta-analysis was not statistically appropriate. [26] The 98 included studies' designs could not rule out selection bias and confounders; thus, the 99 strength of the evidence should be assessed cautiously. Many studies did not adjust for 00 several confounders, which could be responsible for the observed effects. Several studies lacked descriptions of the methods used and the patients included, lowering transparency. 01 02 The results are relevant to health care services, focusing on the well-being and safety of the 23 patients. 04

#### 405 Implications for future research and practice

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	406	This systematic review identified the ambiguity in the PAC interventions offered to the
	407	intervention group. In many studies, it was evident that the methods used lacked clarity, and
	408	high-quality research is needed in this field. The included studies did not demonstrate earlier
	409	surgical room entry time[49,50] or reduction in the number of preoperative tests for
	410	patients attending the PAC, similar to the results of the retrospective studies.[27]
	411	Other implications may include the organisation structure of different PACs and their
	412	functioning. Additionally, the tests that should be part of the assessment at the PACs should
	413	be investigated. The use of technology, such as streaming services, facilitates different
	414	patient groups and might become crucial for reducing human contact and spread of
	415	infection in context of coronavirus disease 2019.
	416	CONCLUSION
1	417	PAC use has reduced the length of stay and surgery cancellation rate at hospitals. However,
	418	the effectiveness of PAC, the major review question, remains unclear, and requires further
	419	research. There is a demand for high-quality studies capturing robust data describing the
	420	quality of care and clinical outcomes for patients requiring anaesthesia. This requires
1	421	increased focus and funding for this specific area of health services research and could,
	422	therefore, lead to implementation of PACs in health care services and improve patient safety
•	423	and perioperative care.
	424	Acknowledgement
1	425	We thank the librarian Ellen Sejerstedt, University of Agder, Kristiansand, for helping with
	426	the search strategy and elimination of duplicate articles.
•	427	Competing Interests
	428	The authors declare that they have no conflict of interest.
	429	Data sharing statement

3 4	430	All data relevant to the study are included within the article or have been uploaded within		
5 6 7	431	supplemental files.		
7 8 9	432	Funding		
10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41	433	This research received no specific grant from any funding agency in the public, commercial		
	434	or not-for-profit sectors.		
	435	Author contributions		
	436	EWK, MF, AO, and TOT: Study design		
	437	EWK, MF, AO: Search and screening of the articles		
	438	EWK: Data extraction		
	439	MF, AO, RCB, and TOT: Control of data extraction		
	440	EWK, MF, AO, and RCB: Quality assessment of the included articles		
	441	EWK: Drafting of the manuscript		
	442	MF, AO, RCB, and TOT: Contribution to and review of the final version of the manuscript		
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2 3 4	596	Figure legends
5 6	597	
7 8 9	598	Figure 1: Preferred Reporting Items for Systematic Reviews and Meta-Analysis (PRISMA).
10 11	599	[23]
12 13 14	600	Figure 2: The Joanna Briggs Institute Critical appraisal checklist for quasi-experimental
15 16	601	studies was used for the risk of bias assessment. [29]
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Figure 2: The Joanna Briggs Institute Critical appraisal checklist for quasiexperimental studies was used for the risk of bias assessment. [30]

# Appendix 1: Search strategies

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## 1. Database Search Strategies

The search mode for CINAHL was Boolean/Phrase. For those searches that are done without search fields - it is automatically searched in the standard fields that CINAHL uses, including words from title, summary, and subject headings. Search 1 and Search 2 are with words in the title (TI in front of the keywords), however this have not been done with the other searches.

Embase and Medline have the same search mode because we are searching for words from title, summary, and subject headings.

### 2. Initial searches, 11 September 2018

#### 2.1. Main Databases

Database: Embase, (Ovid)

Results: n=1287

Search:

1 ((pre-operativ\* or preoperativ\* or pre operativ\*) and (assessment\* or measurement\* or evaluat\*)).ti.

2 (clinic\* or unit\* or nurs\* or outpatient\* or ward\* or center\* or centre\*).ti.

3 1 and 2

4 ((preanaesthe\* or pre-anaesthe\* or pre anaesthe\* or pre-anesthe\* or preanesthe\* or pre anesthe\*) adj4 (assessment\* or measurement\* or evaluat\* or clinic\* or nurs\* or unit\* or outpatient\* or ward\* or center\* or centre\*)).ti,ab. (689)
5 (((pre-admiss\* or preadmiss\*) adj4 (assessment\* or measurement\* or evaluat\*

or clinic\* or unit\* or nurs\* or outpatient\* or ward\* or centre\* or center\*)) and (surg\* or anaesthe\* or anesthe\* or preoperativ\* or pre-operativ\* or preanaesthe\* or pre-anesthe\* or pre-anest

6 ((anaesthe\* or anasthe\* or anesthe\*) adj4 outpatient\* adj4 clinic\*).ti,ab.

7 ((pre-admiss\* or preadmiss\*) adj4 (center\* or centre\*) adj4 (evaluat\* or assessment\* or measurement\*)).ti,ab.

- 8 or/3-7
- 9 limit 8 to yr="1996 -Current"
- 10 limit 9 to (conference abstracts or embase)

#### Database: Medline (Ovid)

Results: n=997

Search:

1 ((pre-operativ\* or preoperativ\* or pre operativ\*) and (assessment\* or measurement\* or evaluat\*)).ti.

2 clinic\* or unit\* or nurs\* or outpatient\* or ward\* or center\* or centre\*).ti.

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15	assessment of measurement )).II,ab.
16	8 or/3-7
17	9 limit 8 to yr="1996 -Current"
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22	Database: Mediline (Ovid)
23	
24	Results: n=1105
25	Search:
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27	measurement* or evaluat*)) ti
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33	clinic* or nurs* or unit* or outpatient* or ward* or center* or centre*)).ti,ab.
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41	7 ((pre-admiss* or preadmiss*) adj4 (center* or centre*) adj4 (evaluat* or
42	assessment* or measurement*)).ti.ab.
43	8 or/3-7
44	0  limit  9  to  yr = "1006  Current"
45	9  mm or  0  yr = 1990 -Current
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47	Database: Cinahl Plus with Full Text (EBSCO host)
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49	Results: n=166
50	Search:
50	1TL ((pre-operativ* or preoperativ* or "pre-operativ*") AND (accessment* or
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58	preanesthe* or "pre anesthe*") N3 (assessment* or measurement* or evaluat* or
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5 ((pre-admiss\* or preadmiss\*) N3 (assessment\* or measurement\* or evaluat\* or clinic\* or unit\* or nurs\* or outpatient\* or ward\* or centre\* or center\*)) AND (surg\* or anaesthe\* or anesthe\* or preoperativ\* or "pre operativ\*" pre-operativ\* or preanaesthe\* or pre-anaesthe\* or pre-anesthe\* or pre-anesthe\* or pre-anesthe\* or pre-anesthe\* or pre-anesthe\* N3 outpatient\* N3 clinic\* 23 (pre-admiss\* or preadmiss\*) N3 (center\* or centre\*) N3 (evaluat\* or assessment\* or measurement\*)

7 S3 OR S4 OR S5 OR S6 OR S7 487

8 S3 OR S4 OR S5 OR S6 OR S7 Limiters - Published Date: 19960101-; Exclude MEDLINE records

# 4. Search update, 4 February 2021

4.1. Main Databases

Database: Embase (Ovid)

Results: n=1572

Search:

1 ((pre-operativ\* or preoperativ\* or pre operativ\*) and (assessment\* or measurement\* or evaluat\*)).ti.

2 (clinic\* or unit\* or nurs\* or outpatient\* or ward\* or center\* or centre\*).ti.

3 1 and 2

4 ((preanaesthe\* or pre-anaesthe\* or pre anaesthe\* or pre-anesthe\* or preanesthe\* or pre anesthe\*) adj4 (assessment\* or measurement\* or evaluat\* or clinic\* or nurs\* or unit\* or outpatient\* or ward\* or center\* or centre\*)).ti,ab.

5 (((pre-admiss\* or preadmiss\*) adj4 (assessment\* or measurement\* or evaluat\* or clinic\* or unit\* or nurs\* or outpatient\* or ward\* or centre\* or center\*)) and (surg\* or anaesthe\* or anesthe\* or preoperativ\* or pre-operativ\* or preanaesthe\* or pre-anaesthe\* or pre-anaesthe\* or pre-anesthe\* or pre-anest

6 ((anaesthe\* or anasthe\* or anesthe\*) adj4 outpatient\* adj4 clinic\*).ti,ab.

7 ((pre-admiss\* or preadmiss\*) adj4 (center\* or centre\*) adj4 (evaluat\* or assessment\* or measurement\*)).ti,ab.

8 or/3-7

9 limit 8 to yr="1996 -Current"

10 limit 9 to (conference abstracts or embase)

Database: Medline (Ovid)

Results: n=1200

Search:

1 ((pre-operativ\* or preoperativ\* or pre operativ\*) and (assessment\* or measurement\* or evaluat\*)).ti.

2 (clinic\* or unit\* or nurs\* or outpatient\* or ward\* or center\* or centre\*).ti.

3 1 and 2

4 ((preanaesthe\* or pre-anaesthe\* or pre anaesthe\* or pre-anesthe\* or preanesthe\* or preanesthe\*) adj4 (assessment\* or measurement\* or evaluat\* or clinic\* or nurs\* or unit\* or outpatient\* or ward\* or center\* or centre\*)).ti,ab.

5 (((pre-admiss\* or preadmiss\*) adj4 (assessment\* or measurement\* or evaluat\* or clinic\* or unit\* or nurs\* or outpatient\* or ward\* or centre\* or center\*)) and (surg\* or anaesthe\* or anesthe\* or preoperativ\* or pre-operativ\* or preanaesthe\* or preanaesthe\* or pre anaesthe\* or pre-anesthe\* or preanesthe\* or pre anesthe\*)).ti,ab. 6 ((anaesthe\* or anasthe\* or anesthe\*) adj4 outpatient\* adj4 clinic\*).ti,ab. 7 ((pre-admiss\* or preadmiss\*) adj4 (center\* or centre\*) adj4 (evaluat\* or assessment\* or measurement\*)).ti,ab. (4) 8 or/3-7 9 limit 8 to yr="1996 -Current" Database: Cinahl Plus with Full Text (EBSCO host) Results: n=209 Search: 1 TI ((pre-operativ\* or preoperativ\* or "pre operativ\*") AND (assessment\* or measurement\* or evaluat\*)) 2 TI (clinic\* or unit\* or nurs\* or outpatient\* or ward\* or center\* or centre\*) 3 S1 AND S2 4 (preanaesthe\* or pre-anaesthe\* or "pre anaesthe\*" or pre-anesthe\* or preanesthe\* or "pre anesthe\*") N3 (assessment\* or measurement\* or evaluat\* or clinic\* or nurs\* or unit\* or outpatient\* or ward\* or centre\* or center\*) 5 ((pre-admiss\* or preadmiss\*) N3 (assessment\* or measurement\* or evaluat\* or clinic\* or unit\* or nurs\* or outpatient\* or ward\* or centre\* or center\*)) AND (surg\* or anaesthe\* or anesthe\* or preoperativ\* or "pre operativ\*" pre-operativ\* or preanaesthe\* or pre-anaesthe\* or pre-anesthe\* or preanesthe\* or "pre anesthe\*") 6 (anaesthe\* or anasthe\* or anesthe\*) N3 outpatient\* N3 clinic\* 30 7 (pre-admiss\* or preadmiss\*) N3 (center\* or centre\*) N3 (evaluat\* or assessment\* or measurement\*) 8 S3 OR S4 OR S5 OR S6 OR S7 9 S3 OR S4 OR S5 OR S6 OR S7 Limiters - Published Date: 19960101-; Exclude MEDLINE records



# PRISMA 2020 Checklist

5 4 5	Section and Topic	ltem #	Checklist item	Location where item is reported
6	TITLE			
7	Title	1	Identify the report as a systematic review.	1
8	ABSTRACT			
9 10	Abstract	2	See the PRISMA 2020 for Abstracts checklist.	2
11	INTRODUCTION	1		
12	Rationale	3	Describe the rationale for the review in the context of existing knowledge.	3-5
13	Objectives	4	Provide an explicit statement of the objective(s) or question(s) the review addresses.	5
14	METHODS			
15	Eligibility criteria	5	Specify the inclusion and exclusion criteria for the review and how studies were grouped for the syntheses.	6
17	Information sources	6	Specify all databases, registers, websites, organisations, reference lists and other sources searched or consulted to identify studies. Specify the date when each source was last searched or consulted.	5-6
18 19	Search strategy	7	Present the full search strategies for all databases, registers and websites, including any filters and limits used.	5-6+App. 1
20 21	Selection process	8	Specify the methods used to decide whether a study met the inclusion criteria of the review, including how many reviewers screened each record and each report retrieved, whether they worked independently, and if applicable, details of automation tools used in the process.	6
22 23 24	Data collection process	9	Specify the methods used to collect data from reports, including how many reviewers collected data from each report, whether they worked independently, any processes for obtaining or confirming data from study investigators, and if applicable, details of automation tools used in the process.	6-7
25 26	Data items	10a	List and define all outcomes for which data were sought. Specify whether all results that were compatible with each outcome domain in each study were sought (e.g. for all measures, time points, analyses), and if not, the methods used to decide which results to collect.	7
27 28		10b	List and define all other variables for which data were sought (e.g. participant and intervention characteristics, funding sources). Describe any assumptions made about any missing or unclear information.	7
29 30	Study risk of bias assessment	11	Specify the methods used to assess risk of bias in the included studies, including details of the tool(s) used, how many reviewers assessed each study and whether they worked independently, and if applicable, details of automation tools used in the process.	6-7
31	Effect measures	12	Specify for each outcome the effect measure(s) (e.g. risk ratio, mean difference) used in the synthesis or presentation of results.	7, 12
32   33	Synthesis methods	13a	Describe the processes used to decide which studies were eligible for each synthesis (e.g. tabulating the study intervention characteristics and comparing against the planned groups for each synthesis (item #5)).	7
34 35 86		13b	Describe any methods required to prepare the data for presentation or synthesis, such as handling of missing summary statistics, or data conversions.	7
37		13c	Describe any methods used to tabulate or visually display results of individual studies and syntheses.	7
38 39		13d	Describe any methods used to synthesize results and provide a rationale for the choice(s). If meta-analysis was performed, describe the model(s), method(s) to identify the presence and extent of statistical heterogeneity, and software package(s) used.	7
40		13e	Describe any methods used to explore possible causes of heterogeneity among study results (e.g. subgroup analysis, meta-regression).	7
41		13f	Describe any sensitivity analyses conducted to assess robustness of the synthesized results.	7
42	Reporting bias assessment	14	Describe any methods used to assess risk of bias due to missing results in a synthesis (arising from reporting biases).	6-7
45	Certainty	15	Describe any methods used to assest coertainty (or contridence) in the doody of evidence for dan out come	6-7

# PRISMA 2020 Checklist

Section and Topic	ltem #	Checklist item	Location where item is reported
assessment			
RESULTS	1		
Study selection	16a	Describe the results of the search and selection process, from the number of records identified in the search to the number of studies included in the review, ideally using a flow diagram.	7
1	16b	Cite studies that might appear to meet the inclusion criteria, but which were excluded, and explain why they were excluded.	7
2 Study 3 characteristics	17	Cite each included study and present its characteristics.	9-10
4 Risk of bias in 5 studies	18	Present assessments of risk of bias for each included study.	12
6 Results of 7 individual studies	19	For all outcomes, present, for each study: (a) summary statistics for each group (where appropriate) and (b) an effect estimate and its precision (e.g. confidence/credible interval), ideally using structured tables or plots.	8-16
8 Results of	20a	For each synthesis, briefly summarise the characteristics and risk of bias among contributing studies.	8-16
9 syntheses 0	20b	Present results of all statistical syntheses conducted. If meta-analysis was done, present for each the summary estimate and its precision (e.g. confidence/credible interval) and measures of statistical heterogeneity. If comparing groups, describe the direction of the effect.	12
1	20c	Present results of all investigations of possible causes of heterogeneity among study results.	12
3	20d	Present results of all sensitivity analyses conducted to assess the robustness of the synthesized results.	12
4 Reporting biases	21	Present assessments of risk of bias due to missing results (arising from reporting biases) for each synthesis assessed.	12
5 Certainty of 6 evidence	22	Present assessments of certainty (or confidence) in the body of evidence for each outcome assessed.	12
7 DISCUSSION			
B Discussion	23a	Provide a general interpretation of the results in the context of other evidence.	16
	23b	Discuss any limitations of the evidence included in the review.	19
1	23c	Discuss any limitations of the review processes used.	19
2	23d	Discuss implications of the results for practice, policy, and future research.	19-20
OTHER INFORMA	TION		
Registration and	24a	Provide registration information for the review, including register name and registration number, or state that the review was not registered.	2-5
6	24b	Indicate where the review protocol can be accessed, or state that a protocol was not prepared.	2-5
7	24c	Describe and explain any amendments to information provided at registration or in the protocol.	2-5
8 Support	25	Describe sources of financial or non-financial support for the review, and the role of the funders or sponsors in the review.	21
Competing interests	26	Declare any competing interests of review authors.	20
Availability of data, code and other materials	27	Report which of the following are publicly available and where they can be found: template data collection forms; data extracted from included studies; data used for all analyses; analytic code; any other materials used in the review.	20-21
5 <i>From:</i> Page MJ, McKe	nzie JE, I	For peer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml Bossuyt PM, Boutron I, Hoffmann TC, Mulrow CD, et al. The PRISMA 2020 statement: an updated guideline for reporting systematic reviews. BMJ 2021;372:n71. doi: 10	.1136/bmj.n71

## PRISMA 2020 Checklist

For more information, visit: http://www.prisma-statement.org/

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