## **Supplementary material**

## Appendix 1 Literature search strategy

All EBM Reviews via Ovid (Cochrane Database of Systematic Reviews (2005+), ACP Journal Club (1991+), Cochrane Central Register of Controlled Trials (CCTR, 1991+), Cochrane Clinical Answers (CCA), Cochrane Methodology Register (2012+), Database of Abstracts of Reviews of Effects (DARE, 2016+), Health Technology Assessment Database (HTA, 2016+), National Health Service Economic Evaluation Databases (NHSEED, 2016+)):

acute-pancreatitis.ab,hw,ti. AND splanchnic.ab,hw,ti. AND (thrombo\* or phlebothrombo\*).ab,hw,ti. AND ((anticoagula\* or anti-coagula\* or anti-thromb\* or anti-thromb\* or heparin\* or LMWH or coumadin\* or warfarin\*).ab,hw,ti. or (anisindione or apolate-sodium or beciparcil or blood-clotting-inhibitor\* or chlorophacinone or coumarin or defibrotide or dextran-sulfate or diphenadione or fluindione or ghilanten or glycerophosphoinositol-inositolphosphodiesterase or glycosaminoglycan-polysulfate or iliparcil or inclacumab or mopidamol or naroparcil or phenindione or tecarfarin or torapsel or tretoquinol or uproleselan).ab,kw,ti.)

Embase via Ovid (1974+): ((acute pancreatitis/or acutepancreatitis.ab,kw,od,ti.) AND (splanchnic blood flow/ or splanchnic.ab,kw,od,ti.) AND (exp vein thrombosis/ or (thrombo\* or phlebothrombo\*).ab,kw,od,ti.) AND (anticoagulation/or exp anticoagulant agent/or (anticoagula\* or anti-coagula\* or antithromb\* or anti-thromb\* or heparin\* or LMWH or coumadin\* or warfarin\*).ab,kw,od,ti,du,tn. or (anisindione or apolate-sodium or beciparcil or blood-clottinginhibitor\* or chlorophacinone or coumarin or defibrotide or dextran-sulfate or diphenadione or fluindione or ghilanten glycerophosphoinositol-inositolphosphodiesterase glycosaminoglycan-polysulfate or iliparcil or inclacumab or mopidamol or naroparcil or phenindione or tecarfarin or torapsel or tretoquinol or uproleselan).ab,du,kw,od,ti,tn.) NOT (exp juvenile/not adult/, exp animal/not exp human/, case report/, case-report.ti.) Limit to English

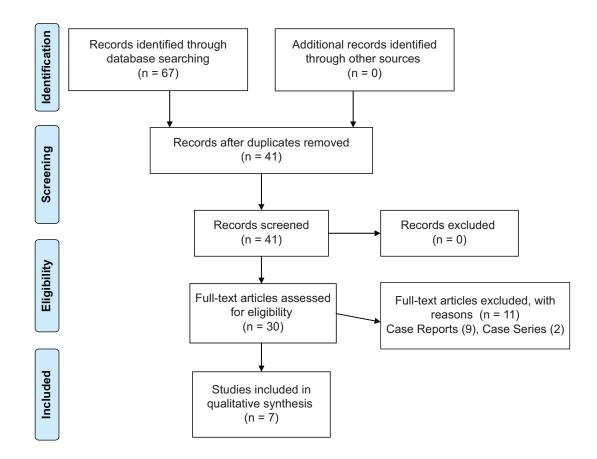
MEDLINE via Ovid (1946+ and Epub Ahead of Print, In-Process & Other Non-Indexed Citations and Ovid MEDLINE(R) Daily): ((exp Pancreatitis/or acute-pancreatitis. ab,kw,ti.) AND (exp Splanchnic Circulation/or splanchnic. ab,kw,ti.) AND (exp Venous Thrombosis/or (thrombo\* or phlebothrombo\*).ab,kw,ti.) AND (exp Anticoagulants/

or (anticoagula\* or anti-coagula\* or antithromb\* or anti-thromb\* or heparin\* or LMWH or coumadin\* or warfarin\*). ab,kw,nm,ti. or (anisindione or apolate-sodium or beciparcil or blood-clotting-inhibitor\* or chlorophacinone or coumarin or defibrotide or dextran-sulfate or diphenadione or fluindione or ghilanten or glycerophosphoinositol-inositolphosphodiesterase or glycosaminoglycan-polysulfate or iliparcil or inclacumab or mopidamol or naroparcil or phenindione or tecarfarin or torapsel or tretoquinol or uproleselan).ab,kw,ti,nm.) NOT (exp CHILD/not exp ADULT/, exp Animals/not Humans/, Case Reports/, case-report.ti.) Limit to English

Scopus via Elsevier (1970+): ((TITLE-ABS-KEY (acute-pancreatitis) AND TITLE-ABS-KEY (splanchnic) AND TITLE-ABS-KEY (thrombo\* OR phlebothrombo\*))) AND (TITLE-ABS-KEY (thrombo\* OR phlebothrombo\*))) AND (TITLE-ABS-KEY (anticoagula\* OR anti-coagula\* OR antithromb\* OR anti-thromb\* OR heparin\* OR lmwh OR coumadin\* OR warfarin\* OR anisindione OR apolate-sodium OR beciparcil OR blood-clotting-inhibitor\* OR chlorophacinone OR coumarin OR defibrotide OR dextransulfate OR diphenadione OR fluindione OR ghilanten OR glycerophosphoinositol-inositolphosphodiesterase OR glycosaminoglycan-polysulfate OR iliparcil OR inclacumab OR mopidamol OR naroparcil OR phenindione OR tecarfarin OR torapsel OR tretoquinol OR uproleselan)) AND (LIMIT-TO (LANGUAGE, "English"))

Web of Science Core Collection via Clarivate Analytics (1975+): TOPIC: (acute-pancreatitis) AND TOPIC: (splanchnic) AND TOPIC: (thrombo\* or phlebothrombo\*) AND TOPIC=(anticoagula\* or anti-coagula\* or antithromb\* or anti-thromb\* or heparin\* or LMWH or coumadin\* or warfarin\* or anisindione or apolate-sodium or beciparcil or blood-clotting-inhibitor\* or chlorophacinone or coumarin or defibrotide or dextran-sulfate or diphenadione or fluindione or ghilanten or glycerophosphoinositol-inositolphosphodiesterase or glycosaminoglycan-polysulfate or iliparcil or inclacumab or mopidamol or naroparcil or phenindione or tecarfarin or torapsel or tretoquinol or uproleselan) Limit to English

Item No	Recommendation	Specify page number										
Reporting of background should include												
1	Problem definition	5										
2	Hypothesis statement	5										
3	Description of study outcome(s)	6										
4	Type of exposure or intervention used	6										
5	Type of study designs used	6										
6	Study population	6-7										
Reporting of search strategy should include												
7	Qualifications of searchers (e.g., librarians and investigators)	6										
8	Search strategy, including time period included in the synthesis and key words	6-7										
9	Effort to include all available studies, including contact with authors	7										
10	Databases and registries searched	6										
11	Search software used, name and version, including special features used (e.g., explosion)	9										
12	Use of hand searching (e.g., reference lists of obtained articles)	5										
13	List of citations located and those excluded, including justification	7, Appendix										
14	Method of addressing articles published in languages other than English	-										
15	Method of handling abstracts and unpublished studies	6										
16	Description of any contact with authors	-										
	Reporting of methods should include											
17	Description of relevance or appropriateness of studies assembled for assessing the hypothesis to be tested	7										
18	Rationale for the selection and coding of data (e.g., sound clinical principles or convenience)	7										
19	Documentation of how data were classified and coded (e.g., multiple raters, blinding and interrater reliability)	7										
20	Assessment of confounding (e.g., comparability of cases and controls in studies where appropriate)	7										
21	Assessment of study quality, including blinding of quality assessors, stratification or regression on possible predictors of study results	7										
22	Assessment of heterogeneity	9										
23	Description of statistical methods (e.g., complete description of fixed or random effects models, justification of whether the chosen models account for predictors of study results, dose-response models, or cumulative meta-analysis) in sufficient detail to be replicated	8-9										
24	Provision of appropriate tables and graphics	Tables 1-2, Figs 1-4										
Reporting of results should include												
25	Graphic summarizing individual study estimates and overall estimate	Figs 1-4										
26	Table giving descriptive information for each study included	Table 1,2										
27	Results of sensitivity testing (e.g., subgroup analysis)	12										
28	Indication of statistical uncertainty of findings	11-12										



## Appendix 3 (B) PRISMA Checklist

Section/topic	#	Checklist item	Specify page number						
		TITLE							
Title	1	Identify the report as a systematic review, meta-analysis, or both.	1						
		ABSTRACT							
Structured summary	2	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.	4						
INTRODUCTION									
Rationale	3	Describe the rationale for the review in the context of what is already known.	5-6						
Objectives	4	Provide an explicit statement of questions being addressed with reference to participants, interventions, comparisons, outcomes, and study design (PICOS).	7-8						
		METHODS							
Protocol and registration	5	Indicate if a review protocol exists, if and where it can be accessed (e.g., Web address), and, if available, provide registration information including registration number.	NA						
Eligibility criteria	6	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 rationale.	7						
Information sources	7	Describe all information sources (e.g., databases with dates of coverage, contact with study authors to identify additional studies) in the search and date last searched.	6						
Search	8	Present full electronic search strategy for at least one database, including any limits used, such that it could be repeated.	6-7 Appendix						
Study selection	9	State the process for selecting studies (i.e., screening, eligibility, included in systematic review, and, if applicable, included in the meta-analysis).	7						
Data collection process	10	Describe method of data extraction from reports (e.g., piloted forms, independently, in duplicate) and any processes for obtaining and confirming data from investigators.	8						
Data items	11	List and define all variables for which data were sought (e.g., PICOS, funding sources) and any assumptions and simplifications made.	8-9						
Risk of bias in individual studies	12	Describe methods used for assessing risk of bias of individual studies (including specification of whether this was done at the study or outcome level), and how this information is to be used in any data synthesis.	12						
Summary measures	13	State the principal summary measures (e.g., risk ratio, difference in means).	8-9						
Synthesis of results	14	Describe the methods of handling data and combining results of studies, if done, including measures of consistency (e.g., $I^2$ ) for each meta-analysis.	10-11						

Supplementary Table 1 Newcastle Ottawa Quality Score

Quality	HIGH>6, MEDIUM 4 to 6, LOW <4	HIGH	HIGH	HIGH	HIGH	HIGH	HIGH	HIGH	
Score	MAX=8	7	7.5		6.5	^	^	7	
Outcome	Adequacy of follow up	All patients followed up: 1; >50% followed up 0.5; <50% followed up OR not mentioned: 0	1	1	1	0.5	1	1	1
	Follow up time	yes: 1; not mentioned: 0	1	1	1	1	1	1	1
	Adequate clinical assessment	yes. 1; no: 0	П	П	П	П	П	П	1
Comparability	Factors comparable between the groups	yes: 1; no: 0	1	1	1	1		п	1
Selection	Outcome not present at start	not present: 1; present: 0	1	1	1	1	1	1	1
	Information on clinical outcomes	Information with clarity: 1; Information derived from percentage value: 0.5; Unclear: 0	1	1	-1	1	П	1	1
	Cohort	>40 patients: 1; 39 to 20: 0.5; <20: 0	1	1	1	1	1	1	1
	Representativeness of the average adult in community	Population based: 1; Multi-center: 0.5; Single-center: 0	0	0.5	0	0	0	0	0
STUDY			Harris, 2013	Junare, 2020	Pagliari, 2020	Toque, 2015	Anderson, 2017 (abs)	Gonzelez, 2011	Garret, 2018