



Universiteit
Leiden
The Netherlands

A meta-analysis of andexanet alfa and prothrombin complex concentrate in the treatment of factor Xa inhibitor-related major bleeding

Jaspers, T.; Shudofsky, K.; Huisman, M.V.; Meijer, K.; Khorsand, N.

Citation

Jaspers, T., Shudofsky, K., Huisman, M. V., Meijer, K., & Khorsand, N. (2021). A meta-analysis of andexanet alfa and prothrombin complex concentrate in the treatment of factor Xa inhibitor-related major bleeding. *Research And Practice In Thrombosis And Haemostasis*, 5(4). doi:10.1002/rth2.12518

Version: Publisher's Version

License: [Creative Commons CC BY-NC-ND 4.0 license](#)

Downloaded from: <https://hdl.handle.net/1887/3230149>

Note: To cite this publication please use the final published version (if applicable).

ORIGINAL ARTICLE

A meta-analysis of andexanet alfa and prothrombin complex concentrate in the treatment of factor Xa inhibitor-related major bleeding

Tessa Jaspers PharmD¹  | Kimberly Shudofsky PharmD² | Menno V. Huisman MD³ | Karina Meijer MD⁴ | Nakisa Khorsand PharmD⁵

¹Department of Clinical Pharmacy and Pharmacology, University Medical Centre Groningen, Groningen, The Netherlands

²Department of Hospital Pharmacy, Viecuri Medical Center, Venlo, The Netherlands

³Department of Thrombosis and Hemostasis, Leiden University Medical Center, Leiden, The Netherlands

⁴Department of Hematology, University Medical Center Groningen, Groningen, The Netherlands

⁵Department of Hospital Pharmacy, OLVG, Amsterdam, The Netherlands

Correspondence

Tessa Jaspers, Department of Clinical Pharmacy and Pharmacology, University Medical Centre Groningen, Hanzeplein 1, 9713 GZ Groningen, The Netherlands.
Email: t.c.c.jaspers@umcg.nl

Funding information

This research did not receive any specific grant from funding agencies in the public, commercial, or not-for-profit sectors.

Handling Editor: Dr Mary Cushman.

Abstract

Background: Andexanet alfa (andexanet) and prothrombin complex concentrate (PCC) are both reversal agents for major bleeding in patients using factor Xa inhibitors (FXaIs). Our aim was to evaluate the current evidence for the effectiveness and safety of andexanet and PCC in a systematic review and meta-analysis.

Objectives: Primary objective was hemostatic effectiveness. Secondary objectives were thromboembolic event rate and mortality.

Methods: A systematic review was performed in PubMed and Embase. Studies describing the effectiveness and/or safety of PCC or andexanet in patients with major bleeding using FXaIs were included. Meta-analysis was performed using a random-effects model.

Results: Seventeen PCC studies, 3 andexanet studies, and 1 study describing PCC and andexanet were included, comprising 1428 PCC-treated and 396 andexanet-treated patients. None of the included studies had control groups, hampering a pooled meta-analysis to compare the two reversal agents. Separate analyses for andexanet and PCC were performed. In subgroup analysis, the pooled proportion of patients with effective hemostasis in studies that used Annexa-4 criteria demonstrated a hemostatic effectiveness of 0.85 (95% confidence interval [CI], 0.80-0.90) in PCC and 0.82 (95% CI, 0.78-0.87) in andexanet studies. The pooled proportion of patients with thromboembolic events was 0.03 (95% CI, 0.02-0.04) in PCC and 0.11 (95% CI, 0.04-0.18) in andexanet studies.

Conclusion: Based on the available evidence with low certainty from observational studies, PCC and andexanet demonstrated a similar, effective hemostasis in the treatment of major bleeding in patients using FXaIs. Compared to PCC, the thromboembolic event rate appeared higher in andexanet-treated patients.

This is an open access article under the terms of the Creative Commons Attribution-NonCommercial-NoDerivs License, which permits use and distribution in any medium, provided the original work is properly cited, the use is non-commercial and no modifications or adaptations are made.

© 2021 The Authors. *Research and Practice in Thrombosis and Haemostasis* published by Wiley Periodicals LLC on behalf of International Society on Thrombosis and Haemostasis (ISTH)

Essentials

- Andexanet alfa and prothrombin complex concentrate (PCC) are both used for major bleeding.
- We evaluated the current evidence for the effectiveness and safety of andexanet alfa and PCC.
- No direct comparisons of both reversal agents were available.
- Both reversal agents demonstrated good hemostatic effectiveness.

1 | INTRODUCTION

Since the introduction of factor Xa inhibitors (FXaIs; rivaroxaban in 2008, apixaban in 2011, and edoxaban in 2015), there has been an urgent need for a reversal agent with proven effectiveness and safety. Although not authorized for this indication, extensive experience has been gained with prothrombin complex concentrate (PCC), both in real life and in clinical trials. Therefore, current guidelines recommend PCC as a therapeutic option in the reversal of FXaI-related major bleeding, based on expert opinion.¹⁻⁵

PCC is known to provide a rapid and complete reversal of vitamin K antagonist (VKA)-induced coagulopathy and is recommended in current guidelines for VKA-associated major bleeding.^{1,5-8} Its safety has been studied extensively. A meta-analysis reported a thromboembolic event rate of 1.9% in patients with VKA coagulopathy treated with PCC.⁹ In 2019, a systematic review and meta-analysis was published concerning the effectiveness and safety of PCC in treating FXaI-related major bleeding.¹⁰ The pooled proportion of patients with hemostatic effectiveness using the ISTH criteria was 0.69 (95% CI, 0.61-0.76), based on two studies involving 150 patients.^{10,11} The pooled proportion of patients with hemostatic effectiveness determined by non-ISTH criteria was 0.77 (95% CI, 0.63-0.92), based on eight studies involving 190 patients.¹⁰

Recently, andexanet alfa (andexanet) was registered by the US Food and Drug Administration (FDA) and European Medicines Agency (EMA) for the indication "neutralization of the anticoagulant activity of apixaban and rivaroxaban in life-threatening or uncontrolled bleeding." This was done by an accelerated procedure based on the registration study because no alternatives for this life-threatening indication were registered.^{12,13} The registration study of andexanet demonstrated a hemostatic effectiveness in 208 of 254 (82%) patients.^{12,13} However, andexanet is not widely available due to its high costs, which form an important obstacle to adding it to the hospital formulary.

Since the registration of andexanet, numerous observational PCC and andexanet studies were published that have further explored their effectiveness and/or safety. Up to now, no meta-analyses are available on the effectiveness and safety of andexanet and PCC for the stated indication. Therefore, the aim of this meta-analysis was to evaluate the current evidence of the effectiveness and safety of andexanet and PCC in the treatment of FXaI-related major bleeding.

2 | METHODS

2.1 | Search strategy

The databases of PubMed and Embase were systematically searched, with articles included up to June 26, 2020. The search strategy can be found in Appendix S1. Inclusion criteria were cohort studies and randomized controlled trials examining the effectiveness and/or safety of andexanet or PCC.

2.2 | Study selection

All studies examining hemostatic effectiveness, thromboembolic events, and/or mortality of andexanet or PCC for the indication major bleeding in patients treated with oral FXaIs were included. As randomized controlled trials on this indication are challenging and therefore could be scarce or absent, observational studies could also be included. Studies with a sample size ≤ 10 patients were excluded, as well as studies from which outcome measures could not be extracted for the subgroup of patients with major bleeding (in studies that also included perioperative use of PCC/andexanet). Two authors (TJ and KS) independently reviewed the title, abstract, and full text of the articles. Disputes were resolved by joint review and consensus. If consensus was not reached, a third author (NK) made the decision whether to include articles.

2.3 | Data extraction

Data of the final set of articles were extracted by TJ and independently checked by KS. Extracted data of interest were study characteristics such as number of patients, study design, inclusion and exclusion criteria, follow-up, and outcomes of interest. For the definition of major bleeding, it was verified whether standardized criteria were used.^{14,15}

2.4 | Risk of bias and study quality assessment

Two authors (TJ and KS) assessed the quality of the observational studies according to two methods. The methodological index for nonrandomized studies (MINORS) method was developed and validated to assess the methodological quality of nonrandomized studies on a 16-point scale.¹⁶

The risk of bias assessment tool for nonrandomized studies (RoBANS) method was developed and validated to determine the bias risk for non-randomized studies.¹⁷ According to the RoBANS method, the bias risk was scored in six different domains: patient selection, risk of confounding, intervention measurement, blinding of assessors, incomplete outcome data, and selective outcome reporting. For each domain, the risk of bias was scored as low, medium, or high. The overall bias risk of each study was classified by showing the mean score as the final score. If the mean score was between two classifications, both classifications were mentioned. Important criteria for the quality assessment were, among others, the inclusion of consecutive patients, usage of standardized criteria to define a major bleeding, prospective collection of data, unbiased evaluation of end points, usage of ISTH criteria to define hemostatic effectiveness, an appropriate follow-up period of ≥ 14 days, and a small loss to follow-up of $\leq 5\%$.

2.5 | Outcome measures

The primary outcome was hemostatic effectiveness as assessed in the included studies. The safety parameters thromboembolic events and mortality were reviewed as secondary outcomes. Thromboembolic events included deep vein thrombosis, pulmonary embolism, myocardial infarction, and/or stroke. Mortality was defined as death from any cause.

2.6 | Meta-analysis

Because no direct comparative studies were found, the safety and effectiveness of andexanet and PCC could not be compared directly by meta-analysis. Therefore, single-arm event rates on hemostatic effectiveness, thromboembolic events and mortality were pooled, using the DerSimonian-Laird random effects model.

Crude pooled proportions with 95% confidence intervals (CIs) of patients with the outcomes were calculated. Forest plots were used to display the results graphically. Subgroup analysis was performed on the used criteria for hemostatic effectiveness: ISTH, Annexa-4, and other nonstandardized criteria. Heterogeneity among studies was assessed with the I^2 statistic. Interpretation of I^2 of 30% to 60% may represent moderate heterogeneity, and substantial heterogeneity was defined as $I^2 > 60\%$.¹⁸ All analyses were performed with Jamovi (version 1.2, Sydney, Australia).

The Preferred Reporting Items for Systematic Reviews and Meta-Analysis guideline was followed. The review protocol of this meta-analysis was not registered prospectively in an international database.

3 | RESULTS

3.1 | Study selection

Using the search criteria, a total of 551 articles were retrieved. After screening, 21 studies were included in the systematic review. The study selection is provided in Figure 1.

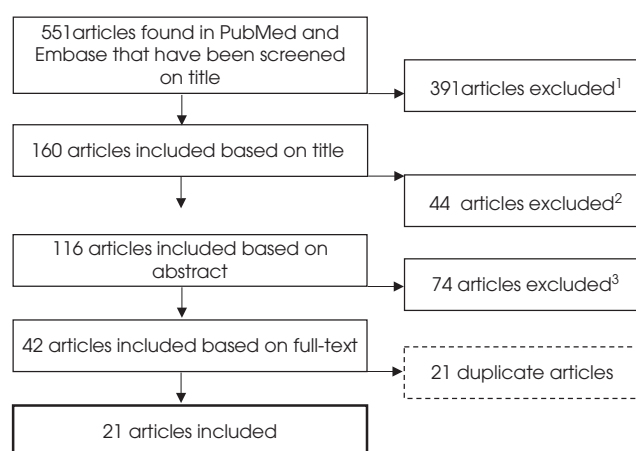


FIGURE 1 Study selection. ¹ Reason for exclusion: updates/reviews: 132, editorial/expert opinion: 50, ex vivo/in vitro study: 44, perioperative usage: 35, no major bleeding in FXa-I users: 30, no PCC or andexanet used: 25, study in healthy volunteers: 19, case reports: 18, guideline: 18, study in animals: 12, cost-efficacy analysis: 4, erratum: 2, survey: 1, preliminary results: 1. ² Reason for exclusion: review: 26, no major bleeding in FXa-I users: 7, ex vivo/in vitro study: 3, healthy volunteers: 2, no relevant outcome measures: 2, preliminary results: 1, no PCC or andexanet used: 1, ≤ 10 patients: 1, survey: 1. ³ Reason for exclusion: Outcome measures could not be extracted for the study population 'major bleeding while using FXa-I, treated with andexanet or PCC': 27, review: 17, editorial/expert opinion: 11, case reports: 6, ≤ 10 patients: 5, no major bleeding in FXa-I users: 5, perioperative usage: 2, in vitro study: 1

3.2 | Study characteristics

Seventeen PCC studies, 3 andexanet studies and 1 study describing PCC and andexanet both were included, comprising 1428 PCC-treated patients and 396 andexanet-treated patients. No direct comparative studies were found, and none of the studies had control groups. Only 4 PCC studies and 1 andexanet study had a prospective design. Of the PCC studies, 7 had a follow-up until discharge, and 10 had a follow-up of ≥ 30 days. All the andexanet studies had a 30-day follow-up. One article (Barra et al³⁵) studied both PCC and andexanet in an observational, retrospective study design with a follow-up of 30 days. Study details are included in Table 1.

Nine PCC studies, one andexanet study, and the study with both reversal agents included only patients with intracerebral hemorrhage (ICH). The vast majority of the included studies had a mean or median age ≥ 75 years. Fifteen PCC studies and two andexanet studies used standardized criteria of the ISTH to define a major bleeding. No other standardized definitions for a major bleeding were used. The registration study of andexanet (Annexa-4), which comprises 352 patients, did not use a standardized definition for a major bleeding.

For hemostatic effectiveness, the standardized ISTH criteria were adopted in five PCC studies (Table 3). In Annexa-4, self-defined criteria to assess hemostatic effectiveness were used. After its publication, three PCC and two other andexanet studies used the Annexa-4 criteria for assessment of hemostatic effectiveness.

TABLE 1 Baseline characteristics of the studies included

Ref.	Treatment	Dose	Study design	No. of FXaI patients	Type of bleeding, %	Type of Fxal, %
Grandhi ¹⁹	PCC	25-50 U/kg	Retrospective, observational, single-center	18	100.0: ICH	88.9: rivaroxaban 11.1: apixaban
Majeed ²⁰	PCC	25 U/kg	Prospective, observational, multicenter	84	70.2: ICH 15.5: GIH 14.3: other	53.6: rivaroxaban 46.4: apixaban
Gerner ²¹	PCC	25-50 U/kg	Retrospective, observational, multicenter	94	100.0: ICH	86.2: rivaroxaban 13.8: apixaban
Schulman ²²	PCC	±25 U/kg (2000 U fixed dose)	Prospective, observational, multicenter	66	55.0: ICH 24.0: GIH 21.0: other	56.1: rivaroxaban 43.9: apixaban
Harrison ²³	PCC	25-50 U/kg	Retrospective, observational, multicenter	14	100.0: ICH	NR
Testa ²⁴	PCC	NR	Prospective, observational, multicenter	27	45.3: ICH 35.9: GIH 18.8: other	74.1: rivaroxaban 25.9: apixaban
Allison ²⁵	PCC	35 U/kg	Retrospective, observational, single-center	33	90.9 ICH 6.1: GIH 3.0 other	81.8: rivaroxaban 18.2: apixaban
Sheikh-Taha (PCC) ²⁶	PCC	50 U/kg	Retrospective, observational, single-center	29	72.4: ICH 13.8: GIH 13.8: other	55.2: rivaroxaban 44.8: apixaban
Arachchillage ²⁷	PCC	25 U/kg	Retrospective, observational, single-center	80	57.5: ICH 30.0: GIH 12.5: other	50.0: rivaroxaban 50.0: apixaban
Stevens ²⁸	Andexanet	PP	Retrospective, observational, single-center	13	46.2: ICH 53.8: other	30.8: rivaroxaban 69.2: apixaban
Dybdahl ²⁹	PCC	50 U/kg	Retrospective, observational, multicenter	35	100.0: ICH	51.4: rivaroxaban 48.6: apixaban
Connolly ¹²	Andexanet	PP, which changed during the study period	Prospective, observational, multicenter	352 for safety analysis 254 for effectiveness analysis	Safety analysis: 64.5: ICH 25.6: GIH 9.9: other Efficacy analysis: 67.3: ICH 24.4: GIH 8.3: other	Safety analysis: 36.4: rivaroxaban 55.1: apixaban 2.8: edoxaban Efficacy analysis: 39.4: rivaroxaban 52.8: apixaban 1.6: edoxaban

Age, y, mean (SD) or median [IQR]	Additional inclusion and exclusion criteria	Definition MBE	Follow-up
79.7 (10.9)	<i>Inclusion:</i> only patients with an ICH <i>Exclusion:</i> use of plasma	ISTH	Until discharge
75.0 [70.0–83.0]	<i>Exclusion:</i> use of other hemostatics (plasma, platelets, aPCC, or factor VIIa), ACS or CVA in the past 30 d, Hb decline without focus, and a do not resuscitate policy	ISTH	30 d
77.5 (7.6)	<i>Inclusion:</i> only patients with an ICH <i>Exclusion:</i> ICH related to trauma, tumor, arteriovenous malformation, aneurysmal subarachnoid hemorrhage, acute thrombolysis, or other coagulopathies	ISTH	3 months
76.9 (10.4)	<i>Exclusion:</i> use of other hemostatics (plasma, platelets, aPCC or factor VIIa), ACS or CVA in past 30 d, Hb decrease without focus and a do not resuscitate policy	ISTH	30 d
74 (8)	<i>Inclusion:</i> only patients with an ICH <i>Exclusion:</i> usage of fresh frozen plasma	ISTH	Unknown
79 [74–85]	NA	ISTH	Until discharge and after six months
73 (14.8)	<i>Exclusion:</i> pregnant and incarcerated patients	A life-threatening or potentially life-threatening bleed requiring emergency surgery or invasive procedure, acute bleeding associated with a fall in Hb ≥ 2 g/dL within 48 h, and bleeding requiring blood product transfusion	Until discharge
73.8 (12.0)	NA	ISTH	Until discharge
76.3 (7.9)	NA	MBE was related to receiving PCC; per protocol, only patients with a MBE (defined by ISTH) were eligible for PCC	30 d
69 (10)	NA	A potentially life-threatening bleeding with signs of hemodynamic instability, bleeding with Hb drop ≥ 2 g/dl (or Hb ≤ 8 g/dl if no baseline), or bleeding in a critical area or organ (retroperitoneal, intra-articular, pericardial, epidural, intracranial, or intramuscular with compartment syndrome)	30 d
78.9 (8.9)	<i>Inclusion:</i> only patients with an ICH	ISTH	Until discharge
Safety analysis: 77.4 (10.8) Efficacy analysis: 77.1 (11.1)	<i>Exclusion:</i> ICH patients with a GCS score ≤ 7 or a hematoma volume of ≥ 60 mL. Patients with scheduled surgery within 12 h of andexanet, life expectancy < 1 mo, thrombotic event in the past 14 d. Patients who have received vitamin K antagonists, dabigatran, PCC, factor VIIa, blood or plasma <i>Additional exclusion criteria for the efficacy analysis:</i> patients with an anti-factor Xa activity < 75 ng/mL or patients without a confirmed MBE	A potentially life-threatening bleeding with signs of hemodynamic instability, bleeding with Hb drop ≥ 2 g/dl (or Hb ≤ 8 g/dl if no baseline), or bleeding in a critical area or organ (retroperitoneal, intra-articular, pericardial, epidural, intracranial, or intramuscular with compartment syndrome)	30 d

(Continues)

TABLE 1 (continued)

Ref.	Treatment	Dose	Study design	No. of FXa patients	Type of bleeding, %	Type of Fxal, %
Smith ³⁰	PCC	25 – 50 U/kg	Retrospective, observational, single-center	31	58.1: ICH 3.2: GIH 38.6: other	45.2: rivaroxaban 54.8: apixaban
Sheikh-Taha (aPCC) ³¹	PCC	25 – 50 U/kg	Retrospective, observational, single-center	35	51.4: ICH 28.6: GIH 20.0: other	42.9: rivaroxaban 57.1: apixaban
Brown ³²	Andexanet	PP	Retrospective, observational, multicenter	13 for safety analysis 11 for effectiveness analysis	100.0: ICH	Safety analysis: 23.1: rivaroxaban 76.9: apixaban
Reynolds ³³	PCC	25 – 50 U/kg	Retrospective, observational, single-center	31 for safety analysis 28 for effectiveness analysis	100.0: ICH	Safety-analysis: 54.8: rivaroxaban 45.2: apixaban
Panos ³⁴	PCC	25 – 50 U/kg	Retrospective, observational, multicenter	663 for safety analysis 433 for effectiveness analysis	100.0: ICH	Safety analysis: 44.8: rivaroxaban 55.2: apixaban Efficacy analysis: 46.0: rivaroxaban 54.0: apixaban
Barra ³⁵	PCC	25 – 50 U/kg	Retrospective, observational, two-arm, single-center	11 PCC	100.0: ICH	72.7: rivaroxaban 27.3: apixaban
	Andexanet	PP		18 andexanet		16.7: rivaroxaban 83.3: apixaban
Bavalia ³⁶	PCC	50 U/kg	Prospective, observational, multicenter	51	59.2: ICH 36.8: GIH 13.2: other	71.0: rivaroxaban 21.1: apixaban 7.9: edoxaban
Korobey ³⁷	PCC	50 U/kg	Retrospective, observational, single-center	59	100.0: ICH	32.2: rivaroxaban 67.8: apixaban
Castillo ³⁸	PCC	25 – 50 U/kg	Retrospective, observational, multicenter	67	100.0: ICH	56.7: rivaroxaban 43.3: apixaban

Note: ISTH, MBE in nonsurgical patients is defined as having a symptomatic presentation and: fatal bleeding; and/or bleeding in a critical area or organ, such as intracranial, intraspinal, intraocular, retroperitoneal, intra-articular, pericardial, or intramuscular with compartment syndrome; and/or bleeding giving a hemoglobin drop of 2 g/dL (1.24 mmol/L) or more, or resulting in transfusion of two of more units of whole blood or red cells.¹⁴ Δ Only patients with ICH are shown; 3 surgical patients and 9 patients of which the severity of the bleed is not described are not shown.

Abbreviations: ACS, acute coronary syndrome; aPCC, activated prothrombin complex concentrate; CVA, cerebrovascular accident; GCS, Glasgow Coma Scale; GIH, gastrointestinal hemorrhage; Hb, hemoglobin; ICH, intracranial hemorrhage; ISTH, International Society on Thrombosis and Haemostasis; IQR, interquartile range; MBE, major bleeding event (uncontrolled or life-threatening bleeding); NA, not applicable; NR, not reported; PCC, prothrombin complex concentrate; PP, per protocol (low-dose 400 mg in 30 min, followed by 480 mg in 2 h, high-dose 800 mg in 30 min, followed by 960 mg in 2 h); SD, standard deviation; TE, thromboembolic event.

3.3 | Study quality

All included studies lacked a control group and therefore had a high risk of confounding and selection bias. The majority of studies had a retrospective design; only four PCC studies and one andexanet study had a prospective patient enrollment. Study quality of observational studies was assessed according to the MINORS and RoBANS methods and are shown in Table 2. Detailed information about the assessment of each included study is included in Appendix S4.

Of the PCC studies, eight were of moderate quality with a low to medium bias risk, and six were of low quality with a medium to medium-high bias risk. The PCC studies by Majeed et al,²⁰ Schulman et al,²² and Bavalia et al³⁶ were high-quality studies with a low risk of bias.

Of the andexanet studies, none were graded as high quality. The andexanet studies by Stevens et al²⁸ and Barra et al³⁵ were graded of moderate quality with a low-medium and medium bias risk, respectively. The studies by Connolly et al¹² and Brown et al³² were graded of low quality with a high bias risk. This was mainly due to strict

Age, y, mean (SD) or median [IQR]	Additional inclusion and exclusion criteria	Definition MBE	Follow-up
74 [69–84]	<i>Exclusion:</i> ICH and GCS score ≤ 7 or ACS or CVA in the past 30 d	Bleed in a critical location, a life-threatening bleed that requires surgery or an invasive procedure, or a bleed that requires a blood transfusion	Until discharge
75.9 (14)	NA	ISTH	Until discharge
Safety analysis: 75.2 (13.5)	NA	ISTH	30 d
Safety analysis: 77 [68–84]	NA	ISTH	7 d
Safety analysis: >65: 14.6% 65–75: 26.4% >75: 59.0% Efficacy analysis: >65: 12.5% 65–75: 24.5% >75: 63.0%	<i>Inclusion:</i> only patients with an ICH <i>Exclusion:</i> pregnant or lactating patients, withdrawal of care within 24 h of admission. Exclusion from effectiveness analysis when a follow-up image was not present within the first 24 h of PCC administration <i>Additional exclusion criteria for the efficacy analysis:</i> patients without a follow-up image of the brain within 24 h of PCC administration	ISTH	Mortality: until discharge TE: 30 d or until discharge
71.0 [68.7–73.2] 83.4 [70.3–88.8]	<i>Inclusion:</i> only patients with an ICH	ISTH	Mortality: until discharge TE: 30 d or until discharge
75 (11)	NA	ISTH	30 d
78.5 (10.9)	<i>Inclusion:</i> only patients with an ICH <i>Exclusion:</i> neuro intervention prior to repeat imaging, died or had care withdrawn prior to repeat imaging	ISTH	Mortality: until discharge TE: 30 d or until discharge
77.6 (13)	<i>Inclusion:</i> only patients with an ICH <i>Exclusion:</i> (acute on) chronic ICH, neurosurgical intervention between baseline and 12-h follow-up scan, or an ICH volume ≥ 60 mL	ISTH	Until discharge

selection criteria, not using an intention-to-treat analysis, and not using standardized criteria for defining major bleeding and assessing hemostatic effectiveness.

pooled outcome proportions are provided in Figure 2. Forest plots of the subgroup analysis for hemostatic effectiveness are provided in Appendix S3.

3.4 | Outcome measures

The study results based on hemostatic effectiveness, thromboembolic events, and mortality are shown in Table 3. Forest plots of the

3.4.1 | Prothrombin complex concentrate studies

Two hundred thirty-three of 1428 PCC-treated patients from safety analyses were excluded from hemostatic effectiveness analyses. In

TABLE 2 The quality of the included studies according to the MINORS and RoBANS methods

Reference	MINORS	RoBANS						
	score	Selection	Confounding	Intervention	Blinding assessor	Incomplete outcome data	Selective reporting	Risk of bias (mean RoBANS score)
Prothrombin complex concentrate studies								
Grandhi ¹⁹	10/16	Medium	Medium	Low	Medium	Medium	Low	Medium
Majeed ²⁰	13/16	Medium	Medium	Low	Low	Low	Low	Low
Gerner ²¹	12/16	Medium	Medium	Medium	Low	Low	High	Medium
Schulman ²²	13/16	Medium	Low	Low	Low	Low	Low	Low
Harrison ²³	6/16	Medium	High	Low	High	High	High	High
Testa ²⁴	12/16	Low	Medium	High	Low	Low	High	Medium
Allison ²⁵	7/16	Medium	Medium	Low	High	Medium	High	Medium
Sheikh-Taha (PCC) ²⁶	7/16	Medium	Medium	Medium	Medium	Medium	Low	Medium
Arachchilage ²⁷	10/16	High	Medium	Low	High	Low	Low	Medium
Dybdahl ²⁹	11/16	Medium	Low	Medium	Medium	Low	Low	Low-Medium
Smith ³⁰	11/16	Medium	High	Low	High	Medium	Low	Medium
Sheikh-Taha (aPCC) ³¹	8/16	Medium	Medium	Medium	Medium	Medium	Low	Medium
Reynolds ³³	9/16	Medium	Low	Low	High	High	High	High
Panos ³⁴	11/16	Medium	Low	Low	Medium	High	High	Medium
Bavalia ³⁶	15/16	Low	Low	Low	Low	Medium	High	Low
Korobey ³⁷	12/16	Medium	Low	Low	Medium	Medium	High	Medium
Castillo ³⁸	8/16	Medium	Low	Low	Medium	High	High	Medium
Andexanet Studies								
Stevens ²⁸	11/16	Medium	Medium	Low	Medium	Low	Low	Low-Medium
Connolly ¹²	9/16	High	High	Medium	High	High	Medium	High
Brown ³²	8/16	High	High	Low	High	Medium	High	High
PCC and andexanet studies								
Barra ³⁵	10/16	High	High	Low	Medium	Medium	Low	Medium

total, hemostatic effectiveness was assessed in 1195 PCC-treated patients. The hemostatic effectiveness ranged from 60.0% to 92.9%. The pooled proportion of patients with good hemostatic effectiveness was 0.77 (95% CI, 0.72-0.82; $I^2=64.8\%$), demonstrating substantial heterogeneity hampering pooled analysis.

Subgroup analysis for hemostatic effectiveness based on the assessment criteria used demonstrated a pooled proportion of 0.69 (95% CI, 0.64-0.75; $I^2 = 0.0\%$) in the three high-quality PCC studies that assessed hemostatic effectiveness by ISTH criteria, 0.85 (95% CI, 0.80-0.90; $I^2 = 41.5\%$) in the three PCC studies that assessed hemostatic effectiveness by Annexa-4 criteria and 0.75 (95% CI, 0.68-0.83; $I^2 = 60.8\%$) in the nine PCC studies that used self-defined criteria for hemostatic effectiveness. There was no heterogeneity between the PCC studies that used ISTH criteria for the definition of hemostatic effectiveness. Moderate heterogeneity was demonstrated between the PCC studies that used Annexa-4 criteria and a substantial heterogeneity between studies that used self-defined criteria.

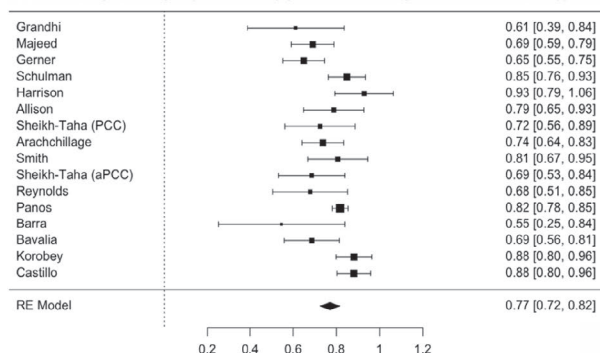
Safety outcome measures were analyzed in 1428 PCC-treated patients. Incidence of thromboembolic events ranged from 0.0% to 12.9%, demonstrating a pooled proportion of 0.03 (95% CI,

0.02-0.04; $I^2 = 18.7\%$). The mortality rate ranged from 13.6% to 63.6%. The pooled analysis on mortality demonstrated substantial heterogeneity and was therefore not further analyzed.

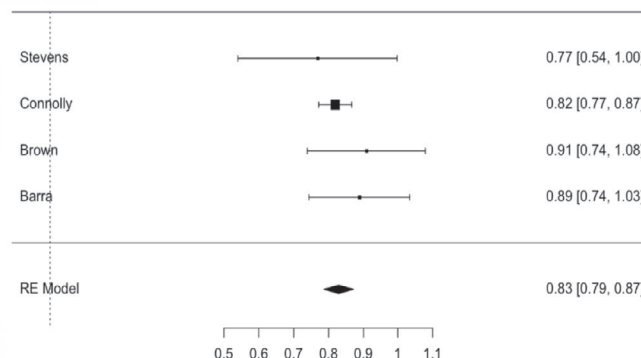
3.4.2 | Andexanet studies

One hundred of 396 andexanet-treated patients from safety analyses were excluded from hemostatic effectiveness analyses. In total, hemostatic effectiveness was assessed in 296 andexanet-treated patients. The hemostatic effectiveness ranged from 77.0% to 90.9%. The pooled proportion of patients with good hemostatic effectiveness was 0.83 (95% CI, 0.79-0.87; $I^2 = 0.0\%$), demonstrating no heterogeneity. Subgroup analysis in the andexanet studies that used Annexa-4 criteria demonstrated a hemostatic effectiveness of 0.82 (95% CI, 0.78-0.87; $I^2 = 0.0\%$). No heterogeneity was observed in the andexanet studies that used Annexa-4 criteria.

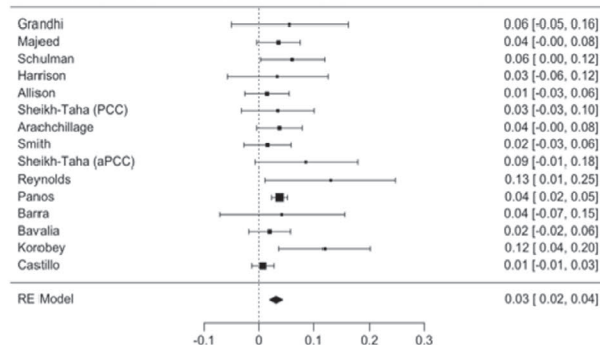
Thromboembolic events and mortality were assessed in 396 andexanet-treated patients. Incidence of thromboembolic events ranged from 0.0% to 30.8% in andexanet studies, demonstrating a pooled proportion of 0.11 (95% CI, 0.04-0.17; $I^2 = 35.7\%$). The

(A) Overall pooled proportions of patients with good hemostatic effectiveness.

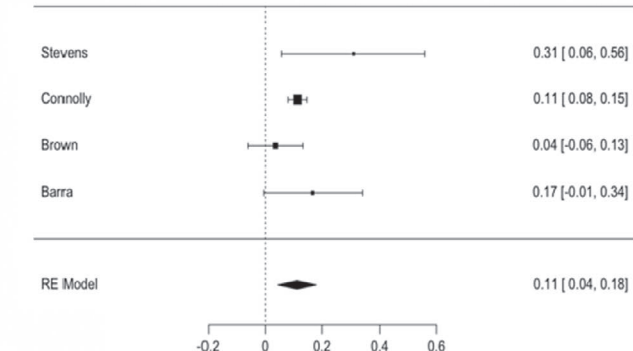
A1. Good hemostatic effectiveness in PCC studies. Pooled proportion is 0.77 (95% CI, 0.72-0.82). Heterogeneity between studies is 64.8%.



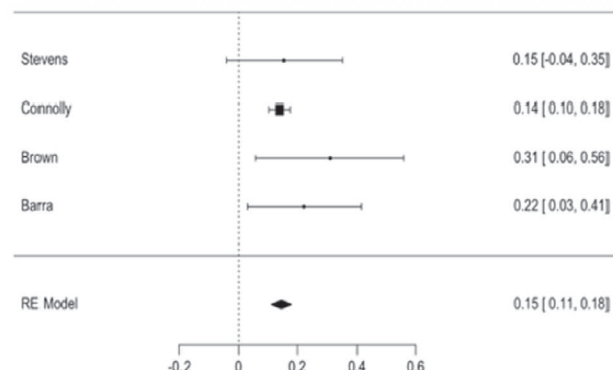
A2. Good hemostatic effectiveness in andexanet studies. Pooled proportion is 0.83 (95% CI, 0.79-0.87). Heterogeneity between studies is 0.0%.

(B) Forest plot of the pooled proportions of patients with thromboembolic events.

B1. Thromboembolic events in PCC studies. The pooled proportion of patients with thromboembolic events is 0.03 (95% CI, 0.02-0.04). Heterogeneity between studies is 18.7%.



B2. Thromboembolic events in andexanet studies. The pooled proportion of patients with thromboembolic events is 0.11 (95% CI, 0.04-0.18). Heterogeneity between studies is 41.1%.

(B) Forest plots of the pooled proportions on mortality.

C1. Mortality in andexanet studies. The pooled proportion is 0.15 (95% CI, 0.11-0.18). Heterogeneity between studies is 0.0%.

FIGURE 2 Forest plots of the pooled outcome proportions. aPCC, activated prothrombin complex concentrate; PCC, prothrombin complex concentrate

mortality rate ranged from 13.9% to 24.0%, demonstrating a pooled proportion of 0.15 (95% CI, 0.11-0.18; $I^2 = 0.0\%$).

4 | DISCUSSION

In this meta-analysis, similar hemostatic effectiveness for PCC and andexanet was observed with pooled proportions of 0.77 and 0.83 in

PCC and andexanet studies, respectively. This was also demonstrated in subgroup analysis in studies that used Annexa-4 criteria for hemostatic effectiveness; a similar hemostatic effectiveness of 0.85 in PCC studies and 0.82 in andexanet studies was demonstrated. The pooled proportion of patients with thromboembolic events was 0.03 (95% CI, 0.02-0.04) in PCC studies and 0.11 (95% CI, 0.04-0.18) in andexanet studies. The mortality rate in PCC studies ranged from 13.6% to 63.6%. The mortality pooled proportion in andexanet studies was 0.15 (95% CI, 0.11-0.18).

TABLE 3 Outcome measures of the included studies

Ref.	Hemostatic effectiveness, n (%)	Criteria for hemostatic effectiveness	Mortality, n (%)	Cause of death and time of onset	Thromboembolic events, n (%)	Definition thromboembolic event and time of onset
PCC studies						
Grandhi ¹⁹	11 (61.1)	No bleeding progression on CT scan	6 (33.3)	4 MBE, 2 pneumonia	1 (5.6)	1 VTE within 24 h
Majeed ²⁰	58 (69.1)	ISTH	27 (32.1)	18 MBE, 7 multiorgan failure, 1 cardiac arrest, 1 PE. 56% 0–7 d, 41% 7–30 d, 3% >30 d	3 (3.6)	2 iCVAs after 5 and 10 d; 1 PE after 15 d
Gerner ²¹	61 (64.9)	No hemorrhagic expansion	NR	NR	NR	NR
Schulman ²²	56 (85.0) 45 (68.0)	Sarode ^a ISTH	9 (13.6)	8 MBE, 1 stab wound. Within 30 d	4 (6.1)	3 iCVAs, 1 DVT, on days 1, 2, 9, 12
Harrison ²³	13 (92.9)	No hemorrhagic expansion	2 (14.3)	NR	0 (0.0)	NA
Testa ²⁴	NR	NR	4 (14.8) ^d	100% within 3 d	NR	NR
Allison ²⁵	26 (83.8)	No bleeding progression on CT scan	5 (15.2)	NR	0 (0.0)	NA
Sheikh-Taha (PCC) ²⁶	21 (72.4)	ISTH	6 (20.7)	6 MBE. 5 within 6 d, 1 on day 14	1 (3.4)	1 iCVA on day 6
Arachchillage ²⁷	59 (73.4)	No recurrent bleeding after 48 h and a patient surviving the MBE	26 (33.0)	NR	3 (3.8)	3 iCVAs within 30 d
Dybdahl ²⁹	NA	NA	8 (22.9)	NR	1 (2.9)	1 VTE
Smith ³⁰	25 (80.6)	Sarode ^a	5 (16.1)	5 MBE	0 (0.0)	NA
Sheikh-Taha (aPCC) ³¹	24 (68.6)	ISTH	7 (20.0)	6 MBE, 1 septic shock, within 9 d	3 (8.6)	2 DVTs on days 2 and 4, 1 MI on day 2
Reynolds ³³	19 (67.9)	Decreased/stable hemorrhage on CT for patients with ICH and ≤20% decrease in Hb 12 h after PCC and no additional blood or factor products within 24 h after PCC for patients without ICH	5 (16.1)	NR	4 (12.9)	3 iCVA, 1 DVT within 7 d
Panos ³⁴	354 (81.8)	Annexa-4 ^b	126 (19.0)	Unknown cause. 48% 0–5 d, 39% 6–14 d, 10% 15–30 d	25 (3.8)	15 DVT, 1 PE, 8 iCVAs, 2 MIs; 50% 0–5 d, 35% 6–14 d, and 15% 15–30 d
Barra ³⁵	6 (60.0)	<35% increase in hematoma volume, SAH thickness, or SDH thickness at 24 h	7 (63.6)	NR	0 (0.0)	NA
Bavalia ³⁶	35 (69.6)	ISTH	9 (17.6)	NR	1 (2.0)	1 PE after 10 d

(Continues)

TABLE 3 (Continued)

Ref.	Hemostatic effectiveness, n (%)	Criteria for hemostatic effectiveness	Mortality, n (%)	Cause of death and time of onset	Thromboembolic events, n (%)	Definition thromboembolic event and time of onset
Korobey ³⁷	52 (88.1)	Annexa-4 ^b	6 (10.2)	NR	7 (11.9)	4 DVTs, 1 PE, 2 iCVAs
Castillo ³⁸	59 (88.1)	Annexa-4 ^b	5 (7.5)	NR	0 (0.0)	NA
Andexanet studies						
Stevens ²⁸	10 (77.0)	Annexa-4 ^b	2 (15.4)	2 MBEs within 3 d	4 (30.8)	1 MI, 1 iCVA, 1 DVT, 1 PE, within 30 d
Connolly ¹²	208 (81.9)	Annexa-4 ^b	49 (13.9)	35 CV cause, 12 non-CV cause, 2 unknown cause.	40 (11.4) ^e	7 MIs, 15 iCVAs, 13 DVTs, 5 PEs; 32% 0-5 d, 32% 6-14 d, 35% 15-30 d
Brown ³²	10 (90.9) ^c	Annexa-4 ^b	4 (30.8) ^f	50% 0-7 d, 50% 7-30 d	0 (0.0) ^f	NA
Barra ³⁵	16 (88.9)	<35% increase in hematoma volume, SAH thickness, or SDH thickness at 24 h	4 (22.2)	NR	3 (16.7)	3 DVTs after 1, 6 and 14 d

Abbreviations: CT, computed tomography; CV, cardiovascular; DVT, deep vein thrombosis; Hb, hemoglobin; Ht, hematocrit; iCVA, ischemic cerebrovascular accident; MI, myocardial infarction; NA, not applicable; NR, not reported; PCC, prothrombin complex concentrate; PE, pulmonary embolism; VTE, venous thromboembolism.

^aThe criteria by Sarode et al were compiled in 2013 with US Food and Drug Administration approval, as there was no standardized method to assess hemostatic effectiveness.³⁷ Its definition is included in Appendix S2

^bThe criteria for effective hemostasis used in Annexa-4 are adjusted Sarode criteria and included in Appendix S2.³⁸

^cPatients who died were excluded from the effectiveness analysis.

^dMortality rate was the same at discharge and at 6 months.

^eForty thromboembolic events occurred in 34 patients.

^fOnly patients with ICH are shown. Three surgical patients and nine patients for which the severity of the bleed is not described are not shown.

All of the included studies lacked a control group, making it difficult to isolate the effect of the reversal agents. A meta-analysis of the registry studies of the direct oral anticoagulants described a case-fatality rate of major bleeding of 7.57% (95% CI, 6.53-8.68).³⁹ This may be underestimated, because patients with comorbidities and concomitant use of antiplatelet agents were excluded in the registration studies. In a prospective study of 732 patients with major bleeding treated with direct oral anticoagulants, the case-fatality rate was 14%.⁴⁰ However, this study does not purely reflect the case-fatality rate without a reversal agent, because 38.4% were treated with PCC.⁴⁰ Nevertheless, the case-fatality rate of 14% was similar to the mortality rates of PCC and andexanet in this meta-analysis. Therefore, it remains uncertain whether PCC and andexanet are of added value in addition to supportive care.

While no difference in hemostatic effectiveness was observed, the incidence of thromboembolic events was higher in the andexanet studies than PCC studies. The pooled proportion of thromboembolic events in PCC-treated patients in this study are consistent with previous studies. Dentali et al⁹ described a thromboembolic event rate of 1.9% in 631 patients who were treated with PCC in VKA-related major bleeding. Piran et al¹⁰ described a 3.0% thromboembolic

event rate in 216 patients who were treated with PCC in FXaI-related major bleeding. Connolly et al¹² stated that the majority of thromboembolic events in Annexa-4 occurred in patients in whom resumption of oral anticoagulation was delayed or in patients who did not restart anticoagulation. However, 32% of the thromboembolic events occurred 0 to 5 days after andexanet administration, and 23.5% occurred after restarting anticoagulation. Furthermore, some hypothesize that andexanet may block tissue factor pathway inhibitor, an endogenous inhibitor of factor Xa, which may lead to thrombosis.⁴¹ A difference in mortality may result in a different ascertained thromboembolic event rate; the thromboembolic event rate is lower in studies with higher mortality, probably due to the competing risk. We conclude that, due to the lack of direct comparative studies, a heterogeneous patient inclusion, and different follow-up periods in included studies, no strong statements can be made about a difference in thromboembolic events between PCC and andexanet treatment.

Our review demonstrates the methodological limitations of the available evidence for effectiveness and safety of PCC and andexanet in FXaI-related major bleeding. All studies were observational without a comparator group, leading to a high risk of selection bias.

Furthermore, the majority of the included studies had retrospective data collection, leading to an even higher risk of bias. Only four PCC studies^{20,22,24,36} minimized the risk of bias through prospective, consecutive patient enrollment, while the only andexanet study with prospective patient enrollment¹² likely suffered a high risk of bias due to strict inclusion criteria, excluding patients with more critical bleeding (eg, patients with a planned intervention within 12 hours after andexanet or an ICH with score <7 on the Glasgow Coma Scale or a hematoma volume >60 mL). Regarding these methodological challenges, we applied two scoring systems to obtain the most objective assessment of the study quality and risk of bias. However, due to the few andexanet studies included, there were insufficient data for a subgroup analysis on moderate/high-quality studies. Eliminating low-quality studies with a high risk of bias would eliminate the studies of Connolly et al¹² and Brown et al,³² eliminating the vast majority of andexanet-treated patients.^{12,32}

We applied several measures to enhance interpretation of the data on hemostatic effectiveness of PCC or andexanet in management of anticoagulation-related major bleeding, such as uniformity in study design characteristics (ISTH criteria for major bleeding) and standardized definitions for hemostatic effectiveness.^{11,14,15} Despite this, heterogeneity among studies was large, hampering pooled analyses.

In the Annexa-4 study, self-defined criteria to assess hemostatic effectiveness were used (included in Appendix S2). The main difference with the ISTH major bleeding criteria is that the assessment time was changed to 12 hours instead of 24 hours after infusion. In addition, no restrictions were made on the administration of hemostatic agents, coagulation factors, or blood products. Moreover, pericardial and intraspinal hemorrhages have their own criteria for effective hemostasis in the Annexa-4 scoring system. These changes may lead to a more favorable outcome of hemostatic effectiveness using the Annexa-4 criteria in comparison to the ISTH criteria.

Annexa-4 is the only study that included anti-factor Xa <75 ng/mL as an exclusion criterion for assessing hemostatic effectiveness. However, FXa levels have short half-lives and may no longer be present in patients with major bleeding, obviating the need for treatment with a reversal agent. Because of this bias, most studies may have overrated the therapy as being effective since anti-factor Xa levels are not usually measured before treatment.

In conclusion, the available evidence makes it challenging to choose a preferred FXa reversal agent. All studies lacked control groups, and the majority were retrospectively conducted with strict selection criteria, without use of standardized definitions for major bleeding, and without standardized scoring methods to define hemostatic effectiveness. Randomized controlled trials with less stringent inclusion criteria and usage of standardized scoring methods are therefore urgently needed. For andexanet, a conditional marketing authorization was given by the EMA and FDA with the condition that a direct comparative study would be conducted to assess the effectiveness and safety of andexanet versus “usual care.” This study, ANNEXA-I, is currently recruiting patients with ICH and is expected to be completed in 2023.⁴²

5 | CONCLUSION

Our systematic review and meta-analysis is the first that we are aware of to evaluate the current evidence for the effectiveness and safety of andexanet and PCC in patients with major bleeding. Separately performed analyses for both reversal agents demonstrated them to be similar in hemostatic effectiveness, while the thromboembolic event rate appeared higher in andexanet. None of the included studies had control groups, hampering a pooled meta-analysis to compare the two reversal agents. There is an urgent need for a randomized clinical trial comparing both reversal agents.

ACKNOWLEDGMENTS

The authors thank B.S. Bhoelan for her support with the meta-analysis.

RELATIONSHIP DISCLOSURE

TJ and KS declare no conflicts of interest. NK has no conflicts of interest for this manuscript. In the past, NK led the ISTH Scientific and Standardization Committee working group, which compiled the ISTH definition for hemostatic effectiveness. MVH has no conflicts of interest for this manuscript. Outside the work, he received grants from ZONMW, Bayer Health Care, Pfizer-BMS, Boehringer-Ingelheim, and Leo Pharma. KM received research grants from Pfizer, Bayer, and Sanquin; lecturing fees from Bayer, Sanquin, Boehringer-Ingelheim, BMS, and Aspen; and consulting fees from Uniqure. All fees were paid to the institute. No other potential conflicts of interest relevant to this article were reported.

AUTHOR CONTRIBUTIONS

TJ contributed to study design, study selection, collection of data, statistical analyses, and writing the initial draft of the manuscript; KS contributed to study design, study selection, and collection of data and provided vital reviews of the manuscript; MVH and KM contributed to study design, interpreted the data, and provided vital reviews of the manuscript. NK contributed to study design and writing the initial draft of the manuscript, interpreted the data, and provided vital reviews of the manuscript.

ORCID

Tessa Jaspers  <https://orcid.org/0000-0002-3939-1403>

REFERENCES

- Christensen H, Cordonnier C, Korv J et al. European stroke organisation guideline on reversal of oral anticoagulants in acute intracerebral haemorrhage. *Eur Stroke J*. 2019;4:294-306.
- Makris M, Van Veen JJ, Tait CR, Mumford AD, Laffan M. British Committee for Standards in Haematology. Guideline on the management of bleeding in patients on antithrombotic agents. *Br J Haematol*. 2013;160:35-46.
- Steffel J, Verhamme P, Potpara TS et al. The 2018 European Heart Rhythm Association Practical Guide on the use of non-vitamin K antagonist oral anticoagulants in patients with atrial fibrillation. *Eur Heart J*. 2018;39:1330-1393.

4. Tomaselli GF, Mahaffey KW, Cuker A et al. 2017 ACC expert consensus decision pathway on management of bleeding in patients on oral anticoagulants: a report of the American College of Cardiology Task Force on Expert Consensus Decision Pathways. *J Am Coll Cardiol*. 2017;70:3042-3067.
5. Witt DM, Nieuwlaat R, Clark NP et al. American Society of Hematology 2018 guidelines for management of venous thromboembolism: optimal management of anticoagulation therapy. *Blood Adv*. 2018;2:3257-3291.
6. Holbrook A, Schulman S, Witt DM et al. Evidence-based management of anticoagulant therapy: antithrombotic therapy and prevention of thrombosis, 9th ed: American College of Chest Physicians Evidence-Based Clinical Practice Guidelines. *Chest*. 2012;141:e152S-e184S.
7. Keeling D, Baglin T, Tait C et al. Guidelines on oral anticoagulation with warfarin - fourth edition. *Br J Haematol*. 2011;154(3):311-324.
8. Khorsand N, Kooistra HA, van Hest RM, Veeger NJ, Meijer K. A systematic review of prothrombin complex concentrate dosing strategies to reverse vitamin K antagonist therapy. *Thromb Res*. 2015;135:9-19.
9. Dentali F, Marchesi C, Giorgi Pierfranceschi M et al. Safety of prothrombin complex concentrates for rapid anticoagulation reversal of vitamin K antagonists. A meta-analysis. *Thromb Haemost*. 2011;106:429-438.
10. Piran S, Khatib R, Schulman S et al. Management of direct factor Xa inhibitor-related major bleeding with prothrombin complex concentrate: a meta-analysis. *Blood Adv*. 2019;3:158-167.
11. Khorsand N, Majeed A, Sarode R et al. Assessment of effectiveness of major bleeding management: proposed definitions for effective hemostasis: communication from the SSC of the ISTH. *J Thromb Haemost*. 2016;14:211-214.
12. Connolly SJ, Crowther M, Eikelboom JW et al. Full study report of andexanet alfa for bleeding associated with factor Xa inhibitors. *N Engl J Med*. 2019;380:1326-1335.
13. U.S. Food & Drug Administration. Accelerated approval. FDA.gov. <https://www.fda.gov/media/113285/download> Published May 3, 2018. Accessed July 1, 2020.
14. Schulman S, Kearon C. Subcommittee on Control of Anticoagulation of the S, Standardization Committee of the International Society on Thrombosis and Haemostasis. Definition of major bleeding in clinical investigations of antihemostatic medicinal products in non-surgical patients. *J Thromb Haemost*. 2005;3:692-694.
15. Palareti G, Leali N, Coccheri S et al. Bleeding complications of oral anticoagulant treatment: an inception-cohort, prospective collaborative study (ISCOAT). Italian Study on Complications of Oral Anticoagulant Therapy. *Lancet*. 1996;348:423-428.
16. Slim K, Nini E, Forestier D, Kwiatkowski F, Panis Y, Chipponi J. Methodological index for non-randomized studies (minors): development and validation of a new instrument. *ANZ J Surg*. 2003;73:712-716.
17. Kim SY, Park JE, Lee YJ et al. Testing a tool for assessing the risk of bias for nonrandomized studies showed moderate reliability and promising validity. *J Clin Epidemiol*. 2013;66:408-414.
18. Higgins JPT, Green S, (editors). Cochrane Handbook for Systematic Reviews of Interventions [Internet] Version 5.1.0 [updated March 2011]. The Cochrane Collaboration, 2011. Chapter 9.5.2, Identifying and measuring heterogeneity. Cited December 1, 2020. Available from: www.handbook.cochrane.org
19. Grandhi R, Newman WC, Zhang X et al. Administration of 4-Factor prothrombin complex concentrate as an antidote for intracranial bleeding in patients taking direct factor Xa inhibitors. *World Neurosurg*. 2015;84:1956-1961.
20. Majeed A, Agren A, Holmstrom M et al. Management of rivaroxaban- or apixaban-associated major bleeding with prothrombin complex concentrates: a cohort study. *Blood*. 2017;130:1706-1712.
21. Gerner ST, Kuramatsu JB, Semblitt JA et al. Association of prothrombin complex concentrate administration and hematoma enlargement in non-vitamin K antagonist oral anticoagulant-related intracerebral hemorrhage. *Ann Neurol*. 2018;83:186-196.
22. Schulman S, Gross PL, Ritchie B et al. Prothrombin complex concentrate for major bleeding on factor Xa inhibitors: a prospective cohort study. *Thromb. Haemost*. 2018;118:842-851.
23. Harrison SK, Garrett JS, Kohman KN, Kline JA. Comparison of outcomes in patients with intracranial hemorrhage on factor Xa inhibitors versus vitamin K antagonists treated with 4-factor prothrombin complex concentrate. *Proc (Bayl Univ Med Cent)*. 2018;31:153-156.
24. Testa S, Ageno W, Antonucci E et al. Management of major bleeding and outcomes in patients treated with direct oral anticoagulants: results from the START-Event registry. *Intern Emerg Med*. 2018;13:1051-1058.
25. Allison TA, Lin PJ, Gass JA et al. Evaluation of the use of low-dose 4-factor prothrombin complex concentrate in the reversal of direct oral anticoagulants in bleeding patients. *J Intensive Care Med*. 2020;35(9):903-908.
26. Sheikh-Taha M. Treatment of apixaban- and rivaroxaban-associated major bleeding using 4-factor prothrombin complex concentrate. *Intern Emerg Med*. 2019;14:265-269.
27. Arachchillage DRJ, Alavian S, Griffin J et al. Effectiveness and safety of prothrombin complex concentrate in patients treated with rivaroxaban or apixaban compared to warfarin presenting with major bleeding. *Br J Haematol*. 2019;184:808-816.
28. Stevens VM, Trujillo T, Mueller SW et al. Coagulation factor Xa (recombinant), inactivated-Zhzo (andexanet alfa) hemostatic outcomes and thrombotic event incidence at an academic medical center. *Clin Appl Thromb Hemost*. 2019;25:1076029619896619.
29. Dybdahl D, Walliser G, Chance Spalding M, Pershing M, Kincaid M. Four-factor prothrombin complex concentrate for the reversal of factor Xa inhibitors for traumatic intracranial hemorrhage. *Am J Emerg Med*. 2019;37:1907-1911.
30. Smith MN, Deloney L, Carter C, Weant KA, Eriksson EA. Safety, effectiveness, and cost of four-factor prothrombin complex concentrate (4F-PCC) in patients with factor Xa inhibitor-related bleeding: a retrospective study. *J Thromb Thrombolysis*. 2019;48:250-255.
31. Sheikh-Taha M, Crawley RM. Reversal of apixaban and rivaroxaban using activated prothrombin complex concentrates in patients with major bleeding. *Am J Cardiovasc Drugs*. 2020;20:295-299.
32. Brown CS, Scott RA, Sridharan M, Rabinstein AA. Real-world utilization of andexanet alfa. *Am J Emerg Med*. 2020;38:810-814.
33. Reynolds TR, Gilbert BW, Hall KM. Utilization of 4-factor prothrombin complex concentrate for reversal of oral factor Xa inhibitor-associated acute major bleeding. *J Pharm Pract*. 2020;897190020907012.
34. Panos NG, Cook AM, John S, Jones GM. Neurocritical Care Society Pharmacy Study Group. Factor Xa inhibitor-related intracranial hemorrhage: results from a multicenter, observational cohort receiving prothrombin complex concentrates. *Circulation*. 2020;141:1681-1689.
35. Barra ME, Das AS, Hayes BD et al. Evaluation of andexanet alfa and four-factor prothrombin complex concentrate (4F-PCC) for reversal of rivaroxaban- and apixaban-associated intracranial hemorrhages. *J Thromb Haemost*. 2020;18:1637-1647.
36. Bavalía R, Abdoellakhan R, Brinkman HJM et al. Emergencies on direct oral anticoagulants: Management, outcomes, and laboratory effects of prothrombin complex concentrate. *Res Pract Thromb Haemost*. 2020;4:569-581.
37. Korobey MJ, Sadaka F, Javed M, Moynihan M, Alsaie A. Efficacy of 4-factor prothrombin complex concentrates in factor Xa inhibitor-associated intracranial bleeding. *Neurocrit Care*. 2021;34(1):112-120.

38. Castillo R, Chan A, Atallah S et al. Treatment of adults with intracranial hemorrhage on apixaban or rivaroxaban with prothrombin complex concentrate products. *J Thromb Thrombolysis*. 2021;51(1):151-158.
39. Chai-Adisaksopha C, Hillis C, Isayama T, Lim W, Iorio A, Crowther M. Mortality outcomes in patients receiving direct oral anticoagulants: a systematic review and meta-analysis of randomized controlled trials. *J Thromb Haemost*. 2015;13:2012-2020.
40. Albaladejo P, Samama CM, Sié P et al. Management of severe bleeding in patients treated with direct oral anticoagulants: an observational registry analysis. *Anesthesiology*. 2017;127:111-120.
41. Levy JH, Ageno W, Chan NC et al. When and how to use antidotes for the reversal of direct oral anticoagulants: guidance from the SSC of the ISTH. *J Thromb Haemost*. 2016;14:623-627.
42. U.S. National Library of Medicine. Trial of Andexanet in ICH Patients Receiving an Oral FXa Inhibitor. Clinicaltrials.gov. <https://clinicaltrials.gov/ct2/show/NCT03661528>. Published September 7, 2018. Updated [July 28, 2020. Accessed August 1, 2020.

SUPPORTING INFORMATION

Additional supporting information may be found online in the Supporting Information section.

How to cite this article: Jaspers T, Shudofsky K, Huisman MV, Meijer K, Khorsand N. A meta-analysis of andexanet alfa and prothrombin complex concentrate in the treatment of factor Xa inhibitor-related major bleeding. *Res Pract Thromb Haemost*. 2021;5:e12518. <https://doi.org/10.1002/rth2.12518>