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Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism (Review)

Robertson L, Jones LE

Robertson L, Jones LE.

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism.

Cochrane Database of Systematic Reviews 2017, Issue 2. Art. No.: CD001100. DOI: 10.1002/14651858.CD001100.pub4.

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thromboembolism (Review)

[Intervention Review]

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

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ABSTRACT

Background

Low molecular weight heparins (LMWHs) have been shown to be effective and safe in preventing venous thromboembolism (VTE). They may also be effective for the initial treatment of VTE. This is the third update of the Cochrane Review first published in 1999.

Objectives

To evaluate the efficacy and safety of fixed dose subcutaneous low molecular weight heparin compared to adjusted dose unfractionated heparin (intravenous or subcutaneous) for the initial treatment of people with venous thromboembolism (acute deep venous thrombosis or pulmonary embolism).

Search methods

For this update the Cochrane Vascular Information Specialist (CIS) searched the Cochrane Vascular Specialised Register (15 September 2016). In addition the CIS searched the Cochrane Central Register of Controlled Trials (CENTRAL; 2016, Issue 8) in the Cochrane Library (searched 15 September 2016) and trials' registries.

Selection criteria

Randomised controlled trials comparing fixed dose subcutaneous LMWH with adjusted dose intravenous or subcutaneous unfractionated heparin (UFH) in people with VTE.

Data collection and analysis

Two review authors independently selected trials for inclusion, assessed for quality and extracted data.

Main results

Six studies were added to this update resulting in a total of 29 included studies (n = 10,390). The quality of the studies was downgraded as there was a risk of bias in some individual studies relating to risk of attrition and reporting bias; in addition several studies did not adequately report on the randomisation methods used nor on how the treatment allocation was concealed.

During the initial treatment period, the incidence of recurrent venous thromboembolic events was lower in participants treated with LMWH than in participants treated with UFH (Peto odds ratio (OR) 0.69, 95% confidence intervals (CI) 0.49 to 0.98; 6238 participants; 18 studies; P = 0.04; moderate-quality evidence). After a follow-up of three months, the period in most of the studies for which oral anticoagulant therapy was given, the incidence of recurrent VTE was lower in participants treated with LMWH than in participants with UFH (Peto OR 0.71, 95% CI 0.56 to 0.90; 6661 participants; 16 studies; P = 0.005; moderate-quality evidence). Furthermore, at the end of follow-up, LMWH was associated with a lower rate of recurrent VTE than UFH (Peto OR 0.72, 95% CI 0.59 to 0.88; 9489 participants; 22 studies; P = 0.001; moderate-quality evidence). LMWH was also associated with a reduction in thrombus size compared to UFH (Peto OR 0.71, 95% CI 0.61 to 0.82; 2909 participants; 16 studies; P < 0.00001; low-quality evidence), but there was moderate heterogeneity (I² = 56%). Major haemorrhages occurred less frequently in participants treated with LMWH than in those treated with UFH (Peto OR 0.69, 95% CI 0.50 to 0.95; 8780 participants; 25 studies; P = 0.02; moderate-quality evidence). There was no difference in overall mortality between participants treated with LMWH and those treated with UFH (Peto OR 0.84, 95% CI 0.70 to 1.01; 9663 participants; 24 studies; P = 0.07; moderate-quality evidence).

Authors' conclusions

This review presents moderate-quality evidence that fixed dose LMWH reduced the incidence of recurrent thrombotic complications and occurrence of major haemorrhage during initial treatment; and low-quality evidence that fixed dose LMWH reduced thrombus size when compared to UFH for the initial treatment of VTE. There was no difference in overall mortality between participants treated with LMWH and those treated with UFH (moderate-quality evidence). The quality of the evidence was assessed using GRADE criteria and downgraded due to concerns over risk of bias in individual trials together with a lack of reporting on the randomisation and concealment of treatment allocation methods used. The quality of the evidence for reduction of thrombus size was further downgraded because of heterogeneity between studies.

PLAIN LANGUAGE SUMMARY

Fixed daily dose of a low molecular weight heparin compared with an adjusted dose of unfractionated heparin for treating blood clots in the deep veins

Background

Venous thromboembolism (VTE) is a condition in which a blood clot forms in the deep veins of the leg or pelvis (DVT) or the clot travels in the blood and blocks a blood vessel in the lungs (pulmonary embolism (PE)). The chances of getting a VTE can be increased if people have risk factors such as previous clots, prolonged periods of immobility (such as travelling on aeroplanes or bed rest), cancer, exposure to oestrogens (pregnancy, oral contraceptives or hormone replacement therapy), trauma and blood disorders such as thrombophilia (abnormal blood clotting). People with a VTE are treated with an anticoagulant, which prevents further clots from forming. Heparin is an anticoagulant and comes in two forms: low molecular weight heparin (LMWH) or unfractionated heparin (UFH). UFH is an older drug and is given either intravenously or by injection. When administering UFH, clinicians have to monitor blood-clotting factors carefully and adjust the dose, because of the variability of its effect. LMWH is given by subcutaneous injection once or twice a day and does not need to be monitored as closely as UFH.

Study characteristics and key results

This review included 29 randomised controlled trials involving 10,390 participants (current to September 2016), which compared LMWH or UFH for treating people with blood clots. Pooling the results of these trials showed that fewer participants treated with LMWH formed further blood clots and that fewer cases of bleeding occurred. Use of LMWH also reduced the size of the original blood clot when compared to the UFH group. There was no difference in number of deaths between participants treated with LMWH and those treated with UFH.

Quality of the evidence

Results of this review indicate that LMWH may prevent further blood clots and bleeding in people with VTE. However, these findings must be interpreted with caution due to the moderate quality of the evidence as a result of lack of reporting of study methods and problems with study design. Results indicating reduced size of blood clots when taking LMWH also must be interpreted with caution due to the low quality of evidence as results were not similar across the studies.

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 2 thromboembolism (Review)

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SUMMARY OF FINDINGS FOR THE MAIN COMPARISON [Explanation]

LMWH compared to UFH for initial treatment of venous thromboembolism

Patient or population: people with venous thromboembolism (VTE) Setting: hospital

Intervention: Low molecular weight heparin (LMWH)

Comparison: Unfractionated heparin (UFH)

Comparison: Uniractionated heparin (UFH)					
Outcomes	Anticipated absolute effects* (95% CI)		Relative effect (95% Cl)	№ of participants (studies)	Quality of the evidence Comments (GRADE)
	Risk with UFH	Risk with LMWH			
Incidence of recurrent	Study population		OR 0.69	6238 (18 RCTs)	⊕⊕⊕⊖ MODERATE ²
VTE ¹ after initial treatment (up to 15 days)	24 per 1000	17 per 1000 (12 to 24)	(0.49 to 0.98)		
Incidence of recurrent	Study population		OR 0.71	6661 (16 RCTs)	⊕⊕⊕⊖ MODERATE ³
VTE ¹ (3 months follow-up)	51 per 1000	37 per 1000 (29 to 46)	(0.56 to 0.90)		
Incidence of recurrent	Study population		OR 0.72 (0.59 to 0.88)	9489 (22 RCTs)	⊕⊕⊕⊖ MODERATE ⁴
VTE ¹ (end of follow-up)	50 per 1000	36 per 1000 (30 to 44)			
Reduction in thrombus	Study population		OR 0.71 (0.61 to 0.82)		⊕⊕⊕⊖ LOW ⁶
size (pre- and post-treat- ment venograms) ⁵	423 per 1000	342 per 1000 (309 to 375)			
Incidence of major haemorrhagic episodes (during initial treatment - up to 15 days) ⁷	Study population		OR 0.69 (0.50 to 0.95)	8780 (25 RCTs)	⊕⊕⊕⊖ MODERATE ⁸

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	21 per 1000	15 per 1000 (11 to 20)			
Overall mortality	Study population		OR 0.84	9663	
(end of follow-up)	57 per 1000	48 per 1000 (41 to 57)	(0.70 to 1.01)	(24 RCTs)	MODERATE ⁹

*The basis for the assumed risk for 'study population' was the average risk in the comparison groups (i.e. total number of participants with events in the control group divided by the number of participants in the comparison group). The risk in the intervention group (and its 95% confidence interval) is based on the assumed risk in the comparison group and the relative effect of the intervention (and its 95% Cl).

CI: Confidence interval; LMWH; low molecular weight heparin; RCTs; randomised controlled trials OR: Peto odds ratio; UFH: unfractionated heparin; VTE: venous thromboembolism

GRADE Working Group grades of evidence

High guality: We are very confident that the true effect lies close to that of the estimate of the effect

Moderate quality: We are moderately confident in the effect estimate: The true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

Low quality: Our confidence in the effect estimate is limited: The true effect may be substantially different from the estimate of the effect

Very low guality: We have very little confidence in the effect estimate: The true effect is likely to be substantially different from the estimate of effect

Recurrent venous thromboembolism (VTE) defined as recurrent deep vein thrombosis (DVT) or recurrent pulmonary embolism (PE). The diagnosis of recurrent DVT was accepted if one of the following criteria was met: (a) a new, constant intraluminal-filling defect not present on the last available venogram; (b) if the venogram was not diagnostic either an abnormal 1251-fibrinogen leg scan or abnormal impedance plethysmogram or ultrasound result, which had been normal before the suspected recurrent episode (Buller 1991). The diagnosis of PE was accepted if one of the following criteria was met: (a) a segmental defect on the perfusion lung scan that was unmatched on the ventilation scan or chest roentgenogram; (b) positive pulmonary angiography; (c) PE at autopsy.

² Downgraded as risk of bias serious due to high risk of attrition bias in 4 studies (Fiessinger 1996; Lindmarker 1994; Ninet 1991; Thery 1992), high risk of reporting bias in 2 studies (Lindmarker 1994; Pérez de Llano 2003) and high risk of other bias in 3 studies (Findik 2002; Harenberg 2000a; Lopaciuk 1992).

³ Downgraded as risk of bias serious due to high risk of attrition bias in 1 study (Breddin 2001), high risk of reporting bias in one study (Pérez de Llano 2003), and high risk of other bias in 2 studies (Findik 2002; Lopaciuk 1992).

⁴ Downgraded as risk of bias serious due to high risk of attrition bias in 2 studies (Breddin 2001; Lindmarker 1994), high risk of reporting bias in 2 studies (Lindmarker 1994; Pérez de Llano 2003), and high risk of other bias in 3 studies (Findik 2002; Harenberg 2000a; Lopaciuk 1992)

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⁵ The number of participants in each group with an improved venographic score, if pre- and post-treatment venograms were obtained and were assessed by persons unaware of treatment assignment.

⁶ Downgraded as risk of bias serious due to high risk of selection bias in 1 study (Luomanmaki 1996), high risk of attrition bias in 6 studies (Breddin 2001; Fiessinger 1996; Kakkar 2003; Lindmarker 1994; Ninet 1991; Thery 1992), high risk of reporting bias in 1 study (Lindmarker 1994), and high risk of other bias in 4 studies (Harenberg 2000a; Kakkar 2003; Lopaciuk 1992; Luomanmaki 1996). Downgraded further due to moderate heterogeneity (I² = 56%)

⁷ Haemorrhages were classified as major if they were intracranial, retroperitoneal, led directly to death, necessitated transfusion or they led to the interruption of antithrombotic treatment or (re)operation.

⁸ Downgraded as risk of bias serious due to high risk of selection bias in 1 study (Luomanmaki 1996), high risk of attrition bias in 5 studies (Fiessinger 1996; Kakkar 2003; Lindmarker 1994; Ninet 1991; Thery 1992), high risk of reporting bias in 2 studies (Lindmarker 1994; Pérez de Llano 2003), and high risk of other bias in 5 studies (Findik 2002; Harenberg 2000a; Kakkar 2003; Lopaciuk 1992; Luomanmaki 1996).

⁹ Downgraded as risk of bias serious due to high risk of selection bias in 1 study (Luomanmaki 1996), high risk of attrition bias in 4 studies (Breddin 2001; Kakkar 2003; Lindmarker 1994; Thery 1992), high risk of reporting bias in 2 studies (Lindmarker 1994; Pérez de Llano 2003), and high risk of other bias in 5 studies (Findik 2002; Harenberg 2000a; Kakkar 2003; Lopaciuk 1992; Luomanmaki 1996).

Fixed dose

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versus adjusted dose unfractionated heparin for the initial treatment of venous

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BACKGROUND

Description of the condition

Venous thromboembolism (presence of a blood clot in the veins, VTE) has an incidence in the general population of approximately 0.1% per year. Its main manifestations are leg complaints, due to deep venous thrombosis (DVT), in the lower limb (blood clot in the deep veins of the leg), and signs of dyspnoea (shortness of breath) and pleuritic thoracic pain (chest pain) when a thrombus (clot) becomes dislodged and forms an embolism obstructing blood flow in the pulmonary circulation. Evidence suggests that although people may only complain about either DVT or pulmonary embolism (PE), in many cases the pathological manifestations are shared between these two clinically distinct conditions (Huisman 1989; Hull 1983). Therefore, increasingly they are referred to as one disease and are treated with comparable anticoagulant regimens.

Description of the intervention

Anticoagulant therapy is the treatment of choice for most people with VTE (NICE 2012). Present guidelines recommend initial therapy for DVT with a parenteral anticoagulant (unfractionated heparin (UFH), low molecular weight heparin (LMWH) or fondaparinux) followed by vitamin K antagonist (VKA) therapy (Kearon 2012). Heparin is administered by either continuous intravenous (IV) infusion or twice daily subcutaneous injection (NICE 2012). Heparin dosage is monitored by the activated partial thromboplastin time (APTT) and adjusted to maintain the anticoagulant effect within a defined therapeutic range. For intravenous heparin therapy to achieve its minimal anticoagulant effect, the initial dosing needs to be either weight based (80 units/kg then 18 units/ kg/hour) or a fixed dose using a 5000 unit bolus followed by at least 1250 units/hour (Kearon 2012). Laboratory monitoring is necessary because the anticoagulant response to heparin is highly variable among people with VTE. Inadequate heparin dosing is related to an increased risk of VTE recurrence (Turpie 2002).

Why it is important to do this review

A number of LMWH preparations and heparinoids have been developed for clinical use. Compared with UFH, LMWH preparations have a longer plasma half-life, less inter-individual variability in anticoagulant response to fixed doses and, in animal models, a more favourable antithrombotic to haemorrhagic ratio (Hirsh 1990; Hirsh 1992). As a result of their pharmacokinetic properties, a stable and sustained anticoagulant effect is achieved when LMWHs are administered subcutaneously once or twice daily, without laboratory monitoring. Although most experience with LMWHs has been in the prevention of VTE, where they have been shown to be safe and effective (Nurmohamed 1992), there is accumulating evidence that these anticoagulants are also safe and effective for the initial treatment of venous thromboembolic events. This is the third update of the Cochrane Review first published in 1999.

OBJECTIVES

To evaluate the efficacy and safety of fixed dose subcutaneous low molecular weight heparin compared to adjusted dose unfractionated heparin (intravenous or subcutaneous) for the initial treatment of people with venous thromboembolism (acute deep venous thrombosis or pulmonary embolism).

METHODS

Criteria for considering studies for this review

Types of studies

Randomised controlled clinical trials (RCTs) with prospective follow-up.

Types of participants

People with venous thromboembolism (acute deep venous thrombosis or pulmonary embolism) confirmed by objective tests.

Types of interventions

Initial treatment (usually in the first five to 14 days) with fixed dose subcutaneous low molecular weight heparin (LMWH) and adjusted dose unfractionated heparin (UFH) (intravenous or subcutaneous).

Types of outcome measures

Primary outcomes

• Incidence of symptomatic recurrent venous thromboembolism (deep venous thrombosis or pulmonary embolism) during the initial treatment and during follow-up.

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Secondary outcomes

• Number of participants in whom the thrombus size reduced based on pre- and post-treatment venograms.

• Frequency of major haemorrhagic episodes during initial treatment or within 48 hours after treatment cessation.

• Overall mortality at the end of follow-up.

Search methods for identification of studies

There were no language restrictions.

Electronic searches

For this update the Cochrane Vascular Information Specialist (CIS) searched the following databases for relevant trials.

• The Cochrane Vascular Specialised Register (searched 15 September 2016).

• The Cochrane Central Register of Controlled Trials

(CENTRAL; 2016, Issue 8) in the Cochrane Library (searched 15 September 2016).

See Appendix 1 for details of the search strategy used to search CENTRAL.

The Cochrane Vascular Specialised Register is maintained by the CIS and is constructed from weekly electronic searches of MED-LINE Ovid, Embase Ovid, CINAHL, AMED, and through handsearching relevant journals. The full list of the databases, journals and conference proceedings which have been searched, as well as the search strategies, used are described in the Specialised Register section of the Cochrane Vascular module in the Cochrane Library (www.cochranelibrary.com).

The CIS searched the following trial registries for details of ongoing and unpublished studies.

• ClinicalTrials.gov (www.ClinicalTrials.gov).

• World Health Organization International Clinical Trials Registry Platform (www.who.int/trialsearch).

• ISRCTN Register (www.isrctn.com/).

Searching other resources

We also reviewed the reference lists of relevant papers identified from these searches.

Data collection and analysis

Selection of studies

For this 2016 update, two review authors (LR and LJ) independently assessed studies identified by the searches for eligibility. Any disagreements were resolved by discussion. Studies were excluded if:

(1) they were dose-ranging studies using higher doses of LMWH than are currently in use;

(2) they used LMWH intravenously;

(3) they adjusted LMWH dosages after initiation of treatment;

(4) the difference in initial treatment was confounded by differences in concomitant medication or long-term medication;

(5) a true LMWH was not used (by true LMWH we mean that no compounds other than heparins were present);

(6) the administration of UFH was suboptimal (i.e. not an adjusted dose);

(7) the report was an abstract with incomplete data.

Data extraction and management

Data were extracted by two review authors (LR and LJ) and included route of administration, intensity of heparin therapy, intensity of oral anticoagulant therapy and the performance of independent assessment of study outcomes.

In addition, the following data were extracted.

(1) The incidence of symptomatic recurrent DVT and PE during the initial treatment and during follow-up (if active follow-up was conducted prospectively at the study centres); whether this incidence was assessed by persons unaware of treatment assignment; and if valid criteria were used for the diagnosis of recurrent VTE. The diagnosis of recurrent DVT was accepted if one of the following criteria was met.

(a) A new, constant intraluminal filling defect not present on the last available venogram.

(b) If the venogram was not diagnostic, either an abnormal 1251fibrinogen leg scan or abnormal impedance plethysmogram or ultrasound result, which had been normal before the suspected recurrent episode (Buller 1991).

The diagnosis of PE was accepted if one of the following criteria was met.

(a) A segmental defect on the perfusion lung scan that was unmatched on the ventilation scan or chest roentgenogram.

(b) Positive pulmonary angiography.

(c) Pulmonary embolism at autopsy.

(2) The number of participants in each group with an improved venographic score, if pre- and post-treatment venograms were obtained and were assessed by persons unaware of treatment assignment.

(3) The frequency of major haemorrhagic episodes during initial treatment. Haemorrhages were classified as major if they were intracranial, retroperitoneal, led directly to death, necessitated transfusion or they led to the interruption of antithrombotic treatment or (re)operation. All other haemorrhages were classified as minor.(4) The overall mortality at the end of follow-up, specified for participants with or without malignant disease, if active follow-up was prospectively conducted at the study centres.

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism (Review)

Assessment of risk of bias in included studies

The risk of bias for all newly included studies was assessed by two review authors (LR and LJ) according to the guidelines given in the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011a). The following domains were assessed as being at either a low risk of bias, high risk of bias or unclear risk of bias using the criteria as described in Chapter 8.5 of the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011b).

1. Sequence generation: was the allocation sequence

adequately generated?

2. Allocation treatment: was allocation adequately concealed?

3. Blinding: was knowledge of the allocated interventions adequately prevented during the study?

4. Incomplete data: were incomplete outcome data adequately addressed?

5. Selective outcome reporting: were reports of the study free of suggestion of selective outcome reporting?

6. Other potential threats to validity: was the study apparently free of other factors that could put it at risk of bias?

We resolved disagreements by discussion and consensus.

Measures of treatment effect

We based reduction in thrombus size on the number of participants whose thrombus size reduced between pre- and post-treatment venograms. We used this outcome and each of the other dichotomous outcomes for the different treatments to calculate an odds ratio (OR) with 95% confidence intervals (CI) separately for each trial. We then combined these ORs across studies, giving due weight to the number of events in each of the two treatment groups in each separate study using the Peto procedure, which assumes a fixed treatment effect (Collins 1987; Mantel 1959). We investigated pulmonary vascular obstruction by calculating the mean difference (MD) between the groups.

We performed all these analyses with the individual LMWH preparations for VTE (that is DVT and PE combined).

We performed an analysis for all LMWH preparations combined if the treatment effects of the individual LMWH preparations were compatible with each other, in view of the biochemical heterogeneity as well as the heterogeneity in animal experiments.

We addressed the validity of combining the trials with a statistical test of homogeneity, which considers whether differences in treatment effect over the individual trials are consistent with natural variation around a constant effect (Collins 1987).

Unit of analysis issues

The unit of analysis in this review was the individual participant.

Dealing with missing data

We sought information about drop-outs, withdrawals and other missing data and, if not reported, we contacted study authors for this information but did not get a response.

Assessment of heterogeneity

We assessed heterogeneity between the trials by visual examination of the forest plot to check for overlapping CIs, the Chi² test for homogeneity with a 10% level of significance and we used the I² statistic to measure the degree of inconsistency between the studies. An I² result of greater than 50% may represent moderate to substantial heterogeneity (Deeks 2011).

Assessment of reporting biases

We assessed publication bias by funnel plots if a sufficient number of studies (10 or more) were available in the meta-analyses. There are many reasons for funnel plot asymmetry, and we consulted the *Cochrane Handbook for Systematic Reviews of Interventions* to aid the interpretation of the results (Sterne 2011).

Data synthesis

One review author (LR) entered the data into Review Manager 5 (RevMan 2014), and the second review author (LJ) cross-checked data entry. We resolved any discrepancies by consulting the source publication. We used a fixed-effect model to meta-analyse the data.

Subgroup analysis and investigation of heterogeneity

We performed subgroup analysis for the different heparin drugs versus unfractionated heparin for all the primary and secondary outcomes of the review.

We also performed the following additional analyses by different groups of interest.

- Proximal deep vein thrombosis.
- Pulmonary embolism.
- Venous thromboembolism with or without malignant disease.
 - Subcutaneous UFH versus LMWH.
 - Intravenous UFH versus LMWH.

For these additional analyses, for the outcome 'recurrent VTE' we report the time point 'end of follow-up' data only.

We also performed a separate analysis to explore any trend over time.

Sensitivity analysis

We performed sensitivity analyses by excluding studies with inadequate concealment of allocation prior to randomisation.

We also performed a sensitivity analysis by excluding studies that did not use the following International Society on Thrombosis and Haemostasis (ISTH) criteria of major bleeding (Schulman 2005).

• Fatal bleeding.

• Symptomatic bleeding in a critical area or organ, such as intracranial, intraspinal, intraocular, retroperitoneal, intra-

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articular or pericardial, or intramuscular with compartment syndrome.

• Bleeding causing a fall in haemoglobin level of 20 g/L (1.24 mmol/L) or more, or leading to transfusion of two or more units of whole blood or red cells.

• Any combination of the above.

Summary of findings

We created 'Summary of findings' tables for LMWH compared with UFH in participants with VTE (Summary of findings for the main comparison). We used GRADEpro GDT software and the GRADE approach to assess the quality of the evidence for the most clinically relevant outcomes as described in Types of outcome measures. We downgraded the evidence from 'high quality' for serious or very serious study limitations (risk of bias, indirectness and inconsistency of evidence, imprecision of effect estimates or potential publication bias) according to the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011a) and the GRADE Working Group (GRADE Working Group 2008).

RESULTS

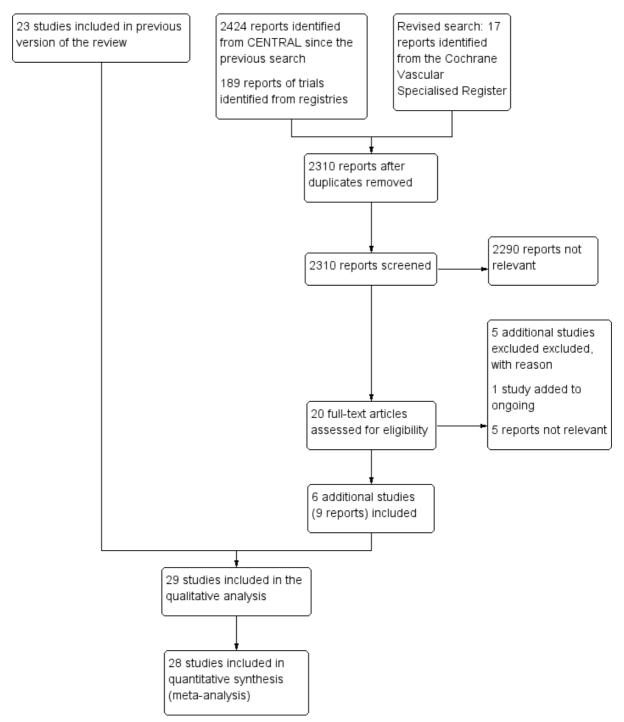
Description of studies

Results of the search

See Figure 1.

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 9 thromboembolism (Review)





Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 10 thromboembolism (Review)

Included studies

Six additional studies were included in this update (Kakkar 2003; Leizorovicz 2011; Meyer 1995; Moreno-Palomares 2001; Pérez de Llano 2003; Thery 1992). In total, 29 studies were truly randomised trials, published between 1988 and the end of 2011, with a total of 10,390 participants. Fourteen of the 29 studies included participants with symptomatic deep venous thrombosis of the leg without symptoms of pulmonary embolism. In eight of these 14 studies people with distal deep venous thrombosis were included as well as people with proximal deep venous thrombosis. In nine studies participants were included if they had symptomatic deep venous thrombosis of the leg, with or without symptomatic pulmonary embolism; or asymptomatic deep venous thrombosis of the leg with symptomatic pulmonary embolism; or symptomatic deep venous thrombosis or pulmonary embolism. In four studies participants with pulmonary embolism only were included. All studies used objective diagnostic tests to confirm the diagnosis. All of the included studies considered fixed dose subcutaneous LMWH once daily (Fiessinger 1996; Hull 1992; Kakkar 2003; Leizorovicz 2011; Lindmarker 1994; Luomanmaki 1996; Simonneau 1997), twice daily (Belcaro 1999; Breddin 2001; Columbus 1997; Decousus 1998; Faivre 1988; Findik 2002; Goldhaber 1998; Harenberg 2000a; Kirchmaier 1998; Koopman 1996; Levine 1996; Lopaciuk 1992; Meyer 1995; Ninet 1991; Pérez de Llano 2003; Prandoni 1992; Prandoni 2004; Riess 2003; Simonneau 1993; Thery 1992), or both (Merli 2001; Moreno-Palomares 2001) compared with adjusted intravenous dose UFH (Breddin 2001; Columbus 1997; Decousus 1998; Fiessinger 1996; Findik 2002; Goldhaber 1998; Harenberg 2000a; Hull 1992; Kakkar 2003; Kirchmaier 1998; Koopman 1996; Levine 1996; Lindmarker 1994; Luomanmaki 1996; Merli 2001; Meyer 1995; Moreno-Palomares 2001; Ninet 1991; Pérez de Llano 2003; Prandoni 1992; Simonneau 1993; Simonneau 1997; Thery 1992) or subcutaneous unfractionated heparin (Faivre 1988; Lopaciuk 1992; Prandoni 2004) or both (Belcaro 1999; Leizorovicz 2011). Nine different preparations of LMWH were identified (nadroparin, tinzaparin, enoxaparin, dalteparin, CY 222, certoparin, ardeparin, reviparin and bemiparin). Ten trials did not have any post-randomisation exclusions or losses to follow-up. Eleven trials reported the number of participants lost to follow-up, which ranged from 1.0% to 12.7%. One trial did not report the dropouts (see Characteristics of included studies).

Excluded studies

Five additional studies were excluded for this update (Quiros 2001; Riess 2014; Siguret 2011; Stricker 1999; Ucar 2015). A total of 26 trials were excluded for the following reasons: dosage of UFH was not adjusted (four trials: Kearon 2006; Notarbartolo 1988; Tedoldi 1993; Zanghi 1988); dose-ranging study (three trials: Banga 1993; de Valk 1995; Handeland 1990); LMWH dosage was adjusted (four trials: Aiach 1989; Bratt 1990; Holm 1986; Ly 1985); intravenous administration of LMWH (four trials: Bratt 1985; Lockner 1985; Lockner 1986; Vogel 1987); results from participants treated for venous thrombosis of the upper limb and for pulmonary embolism could not be distinguished from those of participants with leg vein thrombosis and the outcome was incompletely evaluated (four trials: Albada 1989; Harenberg 1989; Harenberg 1990; Harenberg 2000b); a difference in longterm treatment between the two treatment regimens (two trials: Monreal 1993; Monreal 1994); no UFH comparison group (Siguret 2011); one study looked at the effect of heparin on haemostatic markers and therefore the outcomes were not relevant for this review (Stricker 1999); a substudy of a study already included in the original review (Riess 2003); not an RCT (Quiros 2001); and treatment with thrombolytic therapy (Ucar 2015).

One ongoing study has been identified (NCT00796692). See Characteristics of ongoing studies.

Risk of bias in included studies

See Figure 2.

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous II thromboembolism (Review)



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

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Allocation

Thirteen of the 29 included studies adequately described random sequence generation through the use of a computer or telephone system (Columbus 1997; Decousus 1998; Goldhaber 1998; Hull 1992; Koopman 1996; Leizorovicz 2011; Luomanmaki 1996; Merli 2001; Pérez de Llano 2003; Prandoni 1992; Prandoni 2004; Simonneau 1993; Simonneau 1997). In the remaining 16 studies, there was insufficient information about the random sequence generation to permit a judgement of selection bias. In fourteen of the 29 included studies the assigned treatment was adequately concealed prior to allocation (Columbus 1997; Decousus 1998; Goldhaber 1998; Hull 1992; Kirchmaier 1998; Koopman 1996; Levine 1996; Lopaciuk 1992; Merli 2001; Prandoni 1992; Prandoni 2004; Riess 2003; Simonneau 1993; Simonneau 1997), while in the other 14 trials concealment of allocation was unclear, based on the information given in the publication. One study was deemed to be at high risk of selection bias as there was no central allocation (Luomanmaki 1996). Instead, randomisation was conducted separately at each participating centre (see Characteristics of included studies).

Blinding

In two of the studies, authors did not state whether the participants and staff were blinded to the treatment or not and therefore the risk of performance bias for these two studies was unclear (Harenberg 2000a; Moreno-Palomares 2001). In the remaining 27 included studies treatment allocation was not blinded due to the difference in route of administration between subcutaneous LMWH and intravenous UFH. However, given the clinical outcomes of the study, we judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes and therefore we judged these studies to be at low risk of bias. Even the three studies of subcutaneous UFH versus subcutaneous LMWH were not blinded for treatment allocation due to an initial intravenous bolus in the UFH group (Faivre 1988; Lopaciuk 1992; Prandoni 2004). There was only one double-blinded clinical trial in which participants received either intravenous UFH with subcutaneous placebo or subcutaneous LMWH with intravenous placebo (Hull 1992).

Four of the 29 included studies did not report whether outcome assessors were blinded to treatment and were therefore judged to be at an unclear risk of detection bias (Findik 2002; Goldhaber 1998; Moreno-Palomares 2001; Pérez de Llano 2003). In the remaining 25 studies, outcome assessors were blinded to treatment and therefore these studies were judged to be at low risk of detection bias.

Incomplete outcome data

Six studies were judged to be at high risk of attrition bias as data were missing or imbalanced across the groups (Breddin 2001; Fiessinger 1996; Kakkar 2003; Lindmarker 1994; Ninet 1991; Thery 1992); 18 were judged to be at low risk (Belcaro 1999; Columbus 1997; Decousus 1998; Harenberg 2000a; Hull 1992; Koopman 1996; Leizorovicz 2011; Levine 1996; Lopaciuk 1992; Luomanmaki 1996; Merli 2001; Meyer 1995; Pérez de Llano 2003; Prandoni 1992; Prandoni 2004; Riess 2003; Simonneau 1993; Simonneau 1997); while five did not provide enough information to permit a judgement (Faivre 1988; Findik 2002; Goldhaber 1998; Kirchmaier 1998; Moreno-Palomares 2001).

Selective reporting

Two studies were judged to be at high risk of reporting bias (Lindmarker 1994; Pérez de Llano 2003). In the study by Lindmarker 1994, participants who had died or had a VTE were not included in the analysis. In the study by Pérez de Llano 2003, length of stay was not a prespecified outcome but authors reported data on it in the discussion. Twenty-two studies were at low risk while the remaining five did not provide enough information to permit judgement on reporting bias (Faivre 1988; Fiessinger 1996; Lopaciuk 1992; Moreno-Palomares 2001; Prandoni 1992).

Other potential sources of bias

Five studies were judged to be at high risk of bias (Findik 2002; Harenberg 2000a; Kakkar 2003; Lopaciuk 1992; Luomanmaki 1996). Two studies were sponsored by the pharmaceutical companies that provided the study drug (Harenberg 2000a; Kakkar 2003). The study by Findik 2002 had a low statistical power due to low numbers of participants and few outcome events. Lopaciuk 1992 had an imbalance in exclusion of participants at baseline while the study by Luomanmaki 1996 had a higher incidence of malignancy in participants treated with UFH. Twenty studies were judged to be free from other sources of bias; while in the remaining four, there was not enough information to permit judgement (Decousus 1998; Faivre 1988; Leizorovicz 2011; Moreno-Palomares 2001).

Effects of interventions

See: Summary of findings for the main comparison LMWH compared to UFH for initial treatment of venous thromboembolism

None of the trials individually demonstrated protection from recurrent symptomatic venous thromboembolic complications during the initial treatment period. One trial showed that LMWH

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conferred protection from recurrent symptomatic venous thromboembolic complications at the end of follow-up (Breddin 2001). Only Hull 1992 demonstrated a reduction in major haemorrhage after treatment with LMWH. Six studies showed a reduction in thrombus size, between pre-treatment and post-treatment venograms, in favour of LMWH (Breddin 2001; Goldhaber 1998; Kakkar 2003; Lopaciuk 1992; Prandoni 1992; Simonneau 1993).

Incidence of symptomatic recurrent venous thromboembolism (Analysis 1.1 to Analysis 1.5) ('Summary of findings' table 1)

The occurrence of symptomatic recurrent venous thromboembolism was evaluated during the initial treatment period (Columbus 1997; Decousus 1998; Fiessinger 1996; Findik 2002; Harenberg 2000a; Kakkar 2003; Kirchmaier 1998; Koopman 1996; Levine 1996; Lindmarker 1994; Lopaciuk 1992; Meyer 1995; Ninet 1991; Pérez de Llano 2003; Prandoni 1992; Riess 2003; Simonneau 1993; Simonneau 1997; Thery 1992); at one month's follow-up (Columbus 1997; Levine 1996; Pérez de Llano 2003; Prandoni 1992); at three months' follow-up (Belcaro 1999; Breddin 2001; Columbus 1997; Decousus 1998; Findik 2002; Hull 1992; Koopman 1996; Levine 1996; Lopaciuk 1992; Merli 2001; Meyer 1995; Moreno-Palomares 2001; Pérez de Llano 2003; Prandoni 1992; Prandoni 2004; Simonneau 1993; Simonneau 1997); and at six months' follow-up (Harenberg 2000a; Kirchmaier 1998; Koopman 1996; Lindmarker 1994; Pérez de Llano 2003; Prandoni 1992; Riess 2003). Combining all trials with long-term follow-up gave a comparison of recurrent thromboembolism at the end of follow-up. Although Kakkar 2003 reported incidence of recurrent VTE, there was a query regarding the exact number of participants reported to have this outcome. The author was contacted to clarify the data but did not respond and therefore this study was not included in the analysis.

Analysis of the pooled data from these studies demonstrated a reduction in recurrent venous thromboembolic events with LMWH during the initial treatment period (Peto OR 0.69, 95% CI 0.49 to 0.98; moderate-quality evidence; participants = 6238; studies = 18; P = 0.04); at the end of follow-up (Peto OR 0.72, 95% CI 0.59 to 0.88; participants = 9489; studies = 22; P = 0.0005), at three months' follow-up (Peto OR 0.71, 95% CI 0.56 to 0.90; moderate-quality evidence; participants = 6661; studies = 16; P = 0.005); and at six months' follow-up (Peto OR 0.68, 95% CI 0.48 to 0.96; participants = 2841; studies = 7; P = 0.03). However, at one month's follow-up, no difference was found between LMWH and UFH (Peto OR 0.90, 95% CI 0.56 to 1.44; participants = 1741; studies = 4; P = 0.65).

During the initial treatment, 54 (1.7%) of the 3123 participants allocated to LMWH had thrombotic complications versus 76 (2.4%) of the 3115 participants allocated to UFH. After a follow-up of three months, the period in most of the studies for which oral anticoagulant therapy was given, 122 (3.5%) of the 3440 par-

ticipants treated with LMWH had a recurrent thrombotic event versus 164 (5.2%) of the 3221 participants treated with UFH. When different preparations of heparin were compared, a reduction in recurrent VTE was noted during the initial treatment period for enoxaparin (Peto OR 0.51, 95% CI 0.27 to 0.98; participants = 1143; studies = 5; P = 0.04) and at the end of follow-up for certoparin (Peto OR 0.63, 95% CI 0.40 to 0.99; participants = 2007; 3 studies; P = 0.05) versus UFH. Overall, no differences were observed between the heparin preparations during the initial treatment period and at the end of follow-up.

Reduction in thrombus size (Analysis 1.6)

Venograms were obtained before and after heparin treatment in 16 studies (Breddin 2001; Faivre 1988; Fiessinger 1996; Goldhaber 1998; Harenberg 2000a; Kakkar 2003; Kirchmaier 1998; Lindmarker 1994; Lopaciuk 1992; Luomanmaki 1996; Meyer 1995; Moreno-Palomares 2001; Ninet 1991; Prandoni 1992; Simonneau 1993; Thery 1992). In all studies these venograms were adjudicated by investigators unaware of treatment allocation. The combined results of the 16 studies demonstrated a reduction of thrombus size in 51% of the participants (741 out of 1467) treated with LMWH and in 42% of participants (610 out of 1442) treated with UFH. LMWH was associated with a reduction in thrombus size compared with UFH (Peto OR 0.71, 95% CI 0.61 to 0.82; moderate-quality evidence; participants = 2909; studies = 16; P < 0.00001). However there was moderate heterogeneity in this analysis ($I^2 = 56\%$). When we performed analysis by studies reporting on DVT, the heterogeneity was reduced (I² = 34%) (Analysis 2.4). See also below.

Subgroup analysis showed a difference between the LMWH preparations (P = 0.004). Of the individual LMWH preparations, a better venographic outcome was observed for ardeparin (Peto OR 0.37, 95% CI 0.14 to 0.99), enoxaparin (Peto OR 0.34, 95% CI 0.17 to 0.71), reviparin (Peto OR 0.59, 95% CI 0.43 to 0.80), certoparin (Peto OR 0.70, 95% CI 0.50 to 0.98) and bemiparin (Peto OR 0.42, 95% CI 0.24 to 0.74).

Incidence of major haemorrhage during the initial treatment (Analysis 1.7)

Twenty-five of the included trials evaluated the occurrence of major haemorrhage during the initial treatment (Belcaro 1999; Columbus 1997; Decousus 1998; Faivre 1988; Fiessinger 1996; Findik 2002; Harenberg 2000a; Hull 1992; Kakkar 2003; Kirchmaier 1998; Koopman 1996; Leizorovicz 2011; Levine 1996; Lindmarker 1994; Lopaciuk 1992; Luomanmaki 1996; Meyer 1995; Ninet 1991; Pérez de Llano 2003; Prandoni 1992; Prandoni 2004; Riess 2003; Simonneau 1993; Simonneau 1997; Thery 1992). Analysis of the pooled data showed a reduction in major haemorrhagic complications in favour of LMWH (Peto OR 0.69, 95% CI 0.50 to 0.95; participants = 8780; studies =

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25; moderate-quality evidence; P = 0.02). At the end of the initial treatment period, 65 (1.5%) of the 4333 participants in the LMWH group versus 94 (2.1%) of the 4447 participants in the UFH group suffered a major haemorrhage.

Subgroup analysis showed no difference between the LMWH preparations (P = 0.10).

Overall mortality at the end of follow-up (Analysis 1.8)

Twenty-four studies prospectively evaluated the overall mortality at the end of follow-up (Breddin 2001; Columbus 1997; Decousus 1998; Findik 2002; Goldhaber 1998; Harenberg 2000a; Hull 1992; Kakkar 2003; Kirchmaier 1998; Koopman 1996; Leizorovicz 2011; Levine 1996; Lindmarker 1994; Lopaciuk 1992; Luomanmaki 1996; Merli 2001; Meyer 1995; Pérez de Llano 2003; Prandoni 1992; Prandoni 2004; Riess 2003; Simonneau 1993; Simonneau 1997; Thery 1992). There was no difference in overall mortality at the end of follow-up between participants treated with LMWH and UFH (Peto OR 0.84, 95% CI 0.70 to 1.01; moderate-quality evidence; participants = 9663; studies = 24; P = 0.07). In the LMWH group, 234 (4.7%) of the 5004 participants died versus 265 (5.7%) of the 4659 participants in the UFH group.

When analysed by LMWH preparation, certoparin was the only drug found to be associated with a reduction in overall mortality at the end of follow-up (Peto OR 0.59, 95% CI 0.36 to 0.97; P = 0.04). Overall, no differences were observed between the heparin preparations in mortality at the end of follow-up.

Analysis in participants with proximal deep venous thrombosis (Analysis 2.1 to Analysis 2.6)

A total of 4878 participants with proximal deep venous thrombosis were enrolled in eleven studies (Belcaro 1999; Breddin 2001; Harenberg 2000a; Hull 1992; Kakkar 2003; Koopman 1996; Levine 1996; Moreno-Palomares 2001; Prandoni 1992; Riess 2003; Simonneau 1993). Seven preparations of LMWH were used: nadroparin (three trials, 864 participants), dalteparin (one trial, 30 participants), tinzaparin (one trial, 432 participants), enoxaparin (two trials, 634 participants), reviparin (one trial, 763 participants), certoparin (two trials, 1758 participants) and bemiparin (one trial, 397 participants). In the three-armed trial by Kakkar 2003 two bemiparin groups were compared with an UFH control group. However, in one of the bemiparin groups, participants did not receive concomitant VKA therapy. All other studies included in this review used concomitant VKA therapy and in order for our results to be comparable, data for this group of participants in the Kakkar 2003 study was not included in the analysis.

At the end of follow-up, 80 (3.5%) of the 2303 participants treated with LMWH had a symptomatic recurrent venous thromboembolic event versus 143 (6.0%) of the 2369 participants treated with UFH. This reduction was in favour of LMWH (Peto OR 0.57, 95% CI 0.44 to 0.75; participants = 4672; studies = 10; P < 0.0001) (Analysis 2.1). When analysed by LMWH preparation, reviparin was the only drug associated with a reduction in recurrent VTE (Peto OR 0.31, 95% CI 0.15 to 0.63). Overall, no differences were observed between the heparin preparations in symptomatic recurrent venous thromboembolism at the end of follow-up.

LMWH was also associated with a reduction in the incidence of symptomatic, recurrent deep venous thrombosis as well as a reduction in the incidence of pulmonary embolism (respectively Peto OR 0.61, 95% CI 0.41 to 0.91; participants = 2681; studies = 7; P = 0.02; and Peto OR 0.45, 95% CI 0.28 to 0.74; participants = 3024; studies = 7; P = 0.002) (Analysis 2.2; Analysis 2.3). When analysed by type of LMWH preparation, reviparin and certoparin were the only drugs associated with a reduction in the incidence of pulmonary embolism (respectively Peto OR 0.27, 95% CI 0.10 to 0.73; and Peto OR 0.32, 95% CI 0.11 to 0.92). Overall, no differences were observed between the heparin preparations in symptomatic recurrent deep venous thrombosis and pulmonary embolism at the end of follow-up.

Pooled analysis of two studies demonstrated a reduction of thrombus size in 73% of the participants treated with LMWH and in 56% of participants treated with UFH (Kakkar 2003; Moreno-Palomares 2001). LMWH was associated with a better venographic outcome - Peto OR 0.47, 95% CI 0.27 to 0.80; participants = 230; studies = 2; P = 0.006 (Analysis 2.4) - with the result heavily influenced by the Kakkar 2003 study on bemiparin showing a reduction in thrombus size with LMWH (Peto OR 0.42, 95% CI 0.24 to 0.74; participants = 203; studies = 1; P = 0.003) compared with UFH.

Analysis of the pooled data showed a reduction in major haemorrhagic complications in favour of LMWH (Peto OR 0.50, 95% CI 0.29 to 0.85; participants = 3589; studies = 8; P = 0.01) (Analysis 2.5). At the end of the initial treatment period, 18 (1.0%) of the 1804 participants in the LMWH group versus 37 (2.1%) of the 1785 participants in the UFH group suffered a major haemorrhage. Tinzaparin was the only LMWH preparation associated with reduced rates of major haemorrhagic complications (Peto OR 0.19, 95% CI 0.06 to 0.59). Overall, no differences were observed between the heparin preparations in incidence of major haemorrhages during initial treatment.

Overall mortality at the end of follow-up demonstrated a reduction in favour of LMWH (Peto OR 0.63, 95% CI 0.47 to 0.85; participants = 4331; studies = 9; P = 0.002) (Analysis 2.6). In the LMWH group, 72 (3.3%) of the 2183 participants died versus 112 (5.2%) of the 2148 participants in the UFH group. Certorparin was the only LMWH preparation associated with a reduction in overall mortality (Peto OR 0.54, 95% CI 0.30 to 0.96). Overall, no differences were observed between the heparin preparations in overall mortality at the end of follow-up.

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Analysis in participants with pulmonary embolism (Analysis 3.1)

A total of 1407 participants with pulmonary embolism were enrolled in seven studies (Columbus 1997; Findik 2002; Merli 2001; Meyer 1995; Pérez de Llano 2003; Simonneau 1997; Thery 1992). Four preparations of LMWH were used: tinzaparin (one trial, 612 participants), enoxaparin (three trials, 396 participants), dalteparin (two trials, 128 participants), and reviparin (one trial, 271 participants). In the study by Thery 1992, two other treatment groups were given a high dose of nadroparin (600 and 900 antifactor Xa IU/kg). Data from these groups were not included in the analysis in this review.

All seven studies measured the rate of recurrent thromboembolic events at the end of follow-up. Analysis of pooled data showed no difference between participants treated with LMWH and UFH (Peto OR 0.90, 95% CI 0.50 to 1.61; participants = 1407; studies = 7; P = 0.73) (Analysis 3.1). No individual LMWH preparation was associated with a reduction in the rate of recurrent VTE.

Two studies measured change in thrombus size (Meyer 1995; Thery 1992). Pooled analysis showed no difference in the number of LMWH and UFH participants whose thrombus size improved (Peto OR 1.36, 95% CI 0.23 to 8.16; participants = 106; studies = 2; P = 0.74) (Analysis 3.2). Both studies also measured change in thrombus size according to improvement in the Miller (Thery 1992) or peripheral vascular obstruction score (PVOS) (Meyer 1995). Pooled analysis showed an improvement (MD -3.14, 95%CI -4.39 to -1.90; participants = 106; studies = 2; P < 0.00001) (Analysis 3.3). No individual LMWH preparation was associated with a change in thrombus size.

Three studies measured the incidence of major haemorrhagic complications during initial treatment or within 48 hours after treatment cessation (Meyer 1995; Pérez de Llano 2003; Thery 1992). Pooled analysis showed no difference in the incidence of major bleeding between the LMWH and UFH groups (Peto OR 0.44, 95% CI 0.04 to 4.29; participants = 178; studies = 3; P = 0.48) (Analysis 3.4). However there was significant heterogeneity in this analysis (I² = 58%). No individual LMWH preparation was associated with a reduction in the rate of major haemorrhagic complications.

Three studies measured overall mortality (Meyer 1995; Pérez de Llano 2003; Thery 1992). We found no difference in the overall mortality incidence between the LMWH and UFH groups (Peto OR 1.70, 95% CI 0.17 to 16.71; participants = 178; studies = 3; P = 0.65) (Analysis 3.5). No individual LMWH preparation was associated with reduced overall mortality.

Analysis in participants with venous thromboembolism with or without malignant disease (Analysis 4.1 to Analysis 5.1)

Six studies evaluated mortality at the end of follow-up in participants with and without malignant disease (Columbus 1997; Hull 1992; Lindmarker 1994; Lopaciuk 1992; Prandoni 1992; Simonneau 1997). One of these studies individually showed a reduction in deaths at the end of follow-up with LMWH (Peto OR 0.16, 95% CI 0.03 to 0.72; P = 0.02) (Prandoni 1992). Combining the six studies also demonstrated a reduction in overall mortality in participants with cancer who were treated with LMWH (Peto OR 0.53, 95% CI 0.33 to 0.85; participants = 446; P = 0.009) (Analysis 4.1). In participants without cancer who received LMWH, the reduction in overall mortality of approximately 1% was not different between LMWH and UFH (Peto OR 0.97, 95% CI 0.61 to 1.56; participants = 2139; P = 0.91) (Analysis 5.1). Data on recurrent VTE, reduction in thrombus size and major haemorrhage during initial treatment were not available for the group of participants with or without malignant disease.

Analysis of studies of subcutaneous UFH versus LMWH (Analysis 6.1; Analysis 6.2; Analysis 6.3)

In four studies the UFH in the control group was administered subcutaneously although they did not all report on all outcomes (Faivre 1988; Leizorovicz 2011; Lopaciuk 1992; Prandoni 2004). The analysis of the pooled data from these studies demonstrated no reduction in recurrent venous thromboembolism at the end of follow-up (Peto OR 1.05, 95% CI 0.56 to 1.95; participants = 1403; studies = 3; P = 0.88). However there was significant heterogeneity (I² = 58%). There was no difference in the incidence of major haemorrhagic complications (Peto OR 0.91, 95% CI 0.50 to 1.67; participants = 1471; studies = 4; P = 0.76), nor overall mortality (Peto OR 1.46, 95% CI 0.91 to 2.35; participants = 1403; studies = 3; P = 0.12), between groups treated with subcutaneous UFH and LMWH.

Data on reduction in thrombus size were not available for the group of participants who received subcutaneous UFH versus LMWH.

Analysis of studies of intravenous UFH versus LMWH (Analysis 7.1; Analysis 7.2; Analysis 7.3)

In the 21 studies which compared LMWH with intravenous UFH we found a reduction in recurrent venous thromboembolism at the end of follow-up (Peto OR 0.69, 95% CI 0.56 to 0.86; participants = 8375; studies = 21; P = 0.0007); in major haemorrhages (Peto OR 0.62, 95% CI 0.43 to 0.90; participants = 7309; studies = 21; P = 0.01); and in overall mortality (Peto OR 0.77, 95% CI 0.63 to 0.93; participants = 8260; studies = 21; P = 0.008) (Belcaro 1999; Breddin 2001; Columbus 1997; Decousus 1998; Findik 2002; Goldhaber 1998; Harenberg 2000a; Hull 1992; Kakkar 2003; Kirchmaier 1998; Koopman 1996; Levine 1996; Lindmarker 1994; Merli 2001; Meyer 1995; Pérez de Llano 2003; Prandoni 1992; Riess 2003; Simonneau 1993; Simonneau 1997; Thery 1992).

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Data on reduction in thrombus size were not available for the group of participants who received intravenous UFH versus LMWH.

Sensitivity analysis of studies with adequate concealment of allocation prior to randomisation (Analysis 8.1 to Analysis 8.6)

Fourteen studies had clear concealment of allocation prior to randomisation based on the information given in the publications (Columbus 1997; Decousus 1998; Goldhaber 1998; Hull 1992; Kirchmaier 1998; Koopman 1996; Levine 1996; Lopaciuk 1992; Merli 2001; Prandoni 1992; Prandoni 2004; Riess 2003; Simonneau 1993; Simonneau 1997). The analysis of the pooled data from these studies demonstrated no difference between LMWH and UFH in recurrent venous thromboembolism during the initial treatment period (Peto OR 0.72, 95% CI 0.50 to 1.05; participants = 4862; studies = 10; P = 0.09) nor at three months (Peto OR 0.79, 95% CI 0.60 to 1.02; participants = 5435; studies = 11; P = 0.07). However, LMWH was associated with both a reduction in the incidence of recurrent VTE at the end of followup (Peto OR 0.76, 95% CI 0.60 to 0.96; participants = 6984; studies = 14; P = 0.02) and overall mortality (Peto OR 0.80, 95% CI 0.65 to 0.99; participants = 6984; studies = 14; P = 0.04). Major haemorrhage (Peto OR 0.68, 95% CI 0.45 to 1.03; participants = 6014; studies = 12; P = 0.07) was not different after treatment with LMWH compared with UFH. The reduction in the thrombus size, however, was in favour of LMWH (Peto OR 0.49, 95% CI 0.37 to 0.66; participants = 753; studies = 5; P < 0.00001). Therefore, while reductions in recurrent venous thromboembolism, major haemorrhages and overall mortality were observed in the LMWH group compared with UFH when all studies were combined, in a sensitivity analysis of studies with adequate concealment of treatment allocation before randomisation, no differences were observed in the incidence of recurrent venous thromboembolism during initial treatment and after three months nor in the incidence of major haemorrhages between LMWH and UFH.

Sensitivity analysis of studies that used the International Society on Thrombosis and Haemostasis (ISTH) definition of major and clinically relevant bleeding (Analysis 9.1)

Only one study did not use the ISTH definition of major bleeding and was excluded for the sensitivity analysis (Faivre 1988). Analysis of the pooled data showed a reduction in major haemorrhagic complications in favour of LMWH (Peto OR 0.71, 95% CI 0.52 to 0.98; participants = 8712; studies = 24; P = 0.04). These results are similar to the results from the analysis including all studies irrespective of their definition of major and clinically relevant bleeding (Analysis 1.7).

Trends over time (Analysis 10.1 to Analysis 10.4)

In order to investigate the trend over time, we performed analyses in which all studies were ordered by their date of publication. The forest plots of these analyses did not show an obvious trend over time.

DISCUSSION

Summary of main results

Our review of low molecular weight heparin (LMWH) for the initial treatment of venous thromboembolism (VTE) includes more than 9000 participants and indicates that this drug may be more efficacious than unfractionated heparin (UFH) for preventing recurrent VTE. Many of the included studies reported on other advantages of LMWH over UFH. Firstly, the route of administration (subcutaneous once or twice daily) is more convenient and increases the mobility of participants with VTE. Secondly, the pharmacokinetics are more predictable, which abolishes the need for laboratory monitoring and subsequent dose adjustments. Hence, LMWH can be advocated as the standard therapy for people with confirmed VTE. Treatment in an outpatient setting has been demonstrated to be feasible, safe and cost-effective for people with DVT (Koopman 1996; Levine 1996; van den Belt 1998).

Analysis of all studies, regardless of methodological quality, showed that LMWH was associated with a lower incidence of recurrent VTE at the end of follow-up and at three and six months, with 95% CIs less than one (Peto OR 0.72, 95% CI 0.59 to 0.88, Peto OR 0.71, 95% CI 0.56 to 0.90 and Peto OR 0.68, 95% CI 0.48 to 0.96 respectively) but not after one month follow-up (Peto OR 0.90, 95% CI 0.56 to 1.44). However, when sensitivity analysis was performed on studies that concealed allocation of treatment only, no differences were observed in the incidence of recurrent venous thromboembolism during initial treatment and after three months nor in the incidence of major haemorrhages between LMWH and UFH. We therefore judge that the quality of the evidence is moderate.

When we performed analyses according to the type of VTE index event, the rate of recurrent VTE at the end of follow-up remained lower in DVT participants treated with LMWH compared with DVT participants treated with UFH (Peto OR 0.57, 95% CI 0.44 to 0.75). However, analysis in participants with PE showed no difference in the rate of recurrent VTE between the two treatment groups (Peto OR 0.90, 95% CI 0.50 to 1.61). When we performed analyses according to mode of delivery of UFH, we found that LMWH was associated with fewer recurrent VTEs than intravenous UFH (Peto OR 0.69, 95% CI 0.56 to 0.86) but that there was no difference when LMWH was compared with subcutaneous UFH (Peto OR 1.05, 95% CI 0.56 to 1.95).

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The tendency to improved efficacy with LMWH treatment was not at the cost of a higher rate of major haemorrhage. On the contrary, a reduction in major haemorrhage was demonstrated during the initial treatment period with LMWH. This is largely because the LMWH provides a more stable level of anticoagulation whereas unfractionated heparin dose adjustments may result in more peaks and troughs of anticoagulant effect.

Overall completeness and applicability of evidence

Although these results are promising, there are a number of unresolved issues. Firstly, since only approximately 25% of the participants included in this critical review had a diagnosis of primary pulmonary embolism, it can be argued that more data are required before conclusions can be drawn in this population. Secondly, although the combination of all preparations of LMWH seems logical, and heterogeneity could not be identified, current data do not discriminate between different LMWH preparations. A difference between LMWH preparations was only found for one outcome of the review, reduction in thrombus size. However, studies with large sample sizes and which include comparisons of different preparations are needed to determine whether the efficacy and safety of the individual LMWHs is actually comparable. Thirdly, Prandoni and colleagues noted that the route of administration might be relevant to heparin efficacy (Prandoni 2004). When we limited the analysis to studies that used intravenous UFK, similar results as in the main analyses were observed. When the analysis was confined to those studies that used subcutaneous UFH we found no difference in the incidence of recurrent VTE and major haemorrhages. The lack of difference could be due to the smaller groups in this analysis.

The protocol for this review was published in 1997 and the first version of the review was published in 1998. Initial treatment of VTE has changed since then and, as a result, the current objective of this review is no longer as clinically relevant as before. Therefore, to reflect current practice, future updates of this review will include studies on fixed dose subcutaneous UFH. Additionally, in accordance with current VTE trials on direct-acting oral anticoagulants, future updates will assess symptomatic PE and symptomatic proximal DVT as the primary outcome. We will also assess side effects of treatment other than bleeding as an additional outcome.

Quality of the evidence

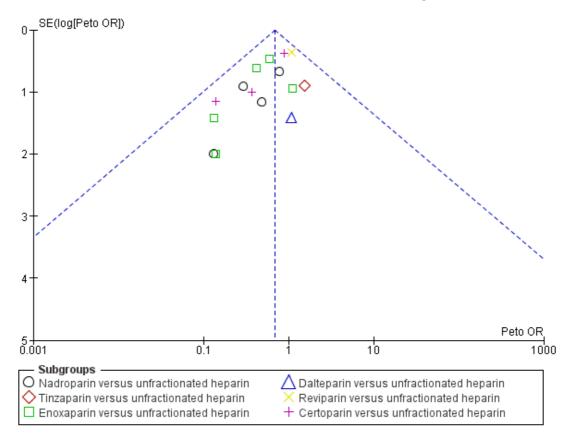
The quality of the evidence was downgraded to moderate due to concerns arising from risk of bias in individual studies. One study was at risk of selection bias (Luomanmaki 1996), six studies were at risk of attrition bias (Breddin 2001; Fiessinger 1996; Kakkar 2003; Lindmarker 1994; Ninet 1991; Thery 1992), two studies were at risk of reporting bias (Lindmarker 1994; Pérez de Llano 2003), and three studies were at risk for other types of bias including baseline differences between the groups (Findik 2002; Lopaciuk 1992; Luomanmaki 1996). A further reason for downgrading the evidence to moderate was that several studies did not adequately report on the methods used to generate the random sequence nor how treatment allocation was kept concealed.

While reductions in recurrent VTE and major haemorrhages were observed in the LMWH group compared with UFH when all studies were combined, in a sensitivity analysis of studies with adequate concealment of treatment allocation before randomisation, no differences were observed in the incidence of recurrent VTE during initial treatment and after three months nor in the incidence of major haemorrhages between LMWH and UFH. An explanation for these differences in effect size could be that the overall reductions are possibly biased by including less adequately performed studies without adequate concealment.

Where there were 10 or more studies in an analysis, we tested for publication bias using funnel plots. We found a suggestion of publication bias for three of the outcomes: incidence of recurrent VTE during initial treatment (Analysis 1.1, Figure 3);incidence of recurrent VTE at three months (Analysis 1.4, Figure 4); and reduction in thrombus size (Analysis 1.6, Figure 5). However, we felt it was insufficient to downgrade for publication bias. For the remaining outcomes, we found no evidence of publication bias for the analyses we tested (Analysis 1.2; Analysis 1.7; Analysis 1.8; Analysis 2.1; Analysis 7.1; Analysis 7.2; Analysis 7.3; Analysis 8.1; Analysis 8.2; Analysis 8.3; Analysis 8.5; Analysis 8.6; Analysis 10.1; Analysis 10.2; Analysis 10.3; Analysis 10.4).

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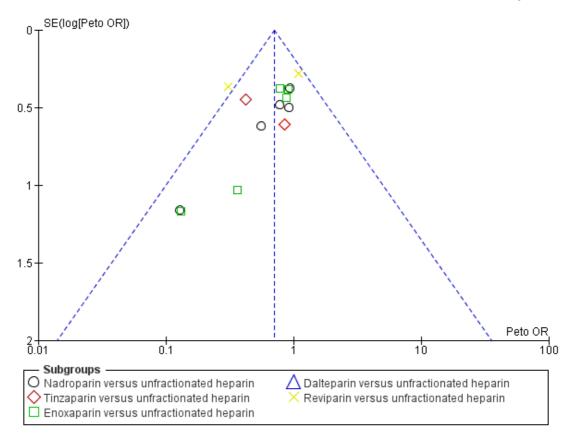
Figure 3. Funnel plot of comparison: I LMWH versus UFH in people with venous thromboembolism, outcome: 1.1 Incidence of recurrent venous thromboembolism during initial treatment.



Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 19 thromboembolism (Review)

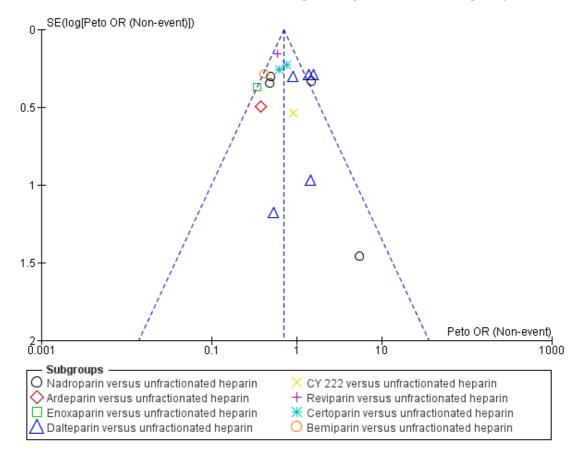
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Figure 4. Funnel plot of comparison: I LMWH versus UFH in people with venous thromboembolism, outcome: 1.4 Incidence of recurrent venous thromboembolism at 3 months' follow-up.



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Figure 5. Funnel plot of comparison: I LMWH versus UFH in people with venous thromboembolism, outcome: 1.6 Reduction in thrombus size (pre- and post-treatment venograms).



Potential biases in the review process

Neither of the authors of this review was involved in any of the included or excluded studies. Furthermore, neither has any commercial or other conflict of interest. The search was as comprehensive as possible; and the two review authors independently assessed all studies for inclusion. We are confident that we have included all relevant studies and we have attempted to reduce bias in the review process by performing data extraction and assessing study quality independently. However, the possibility remains that we may have missed studies which have not been published.

The original review did not set out to use the ISTH bleeding definition. However, given that this is now the standard accepted definition for major bleeding, we performed a post hoc sensitivity analysis for ISTH bleeding definitions in order to assess the effect of bleeding definitions used. The results from this sensitivity analysis (Analysis 9.1) are similar to the results from the analysis including all studies (Analysis 1.7) irrespective of their definition

of major and clinically relevant bleeding.

Agreements and disagreements with other studies or reviews

One network meta-analysis of four studies compared three LMWH preparations (tinzaparin, nadroparin and enoxaparin) in terms of safety and efficacy for the treatment of deep vein thrombosis (Diaz 2015). Authors found no evidence of differences between tinzaparin, nadroparin and enoxaparin for recurrence of DVT and major bleeding.

AUTHORS' CONCLUSIONS

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Implications for practice

This review presents moderate-quality evidence that fixed dose LMWH reduced the incidence of recurrent thrombotic complications and occurrence of major haemorrhage during initial treatment and low-quality evidence that fixed dose LMWH reduced thrombus size when compared to UFH for the initial treatment of VTE. There was no difference in overall mortality between participants treated with LMWH and those treated with UFH (moderate-quality evidence).

Implications for research

Further studies are required to compare LMWH with UFH in the treatment of people with pulmonary embolism. In addition, a large RCT of at least two years' duration should be performed to determine the effects of dosing frequency on long-term sequelae of venous thromboembolism, such as the development of postthrombotic syndrome. Individual low molecular weight heparin preparations could be compared with each other and new drugs should now be compared with LMWH.

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* Indicates the major publication for the study

CHARACTERISTICS OF STUDIES

Characteristics of included studies [ordered by study ID]

Belcaro 1999

Methods	Study design: randomised controlled trial. Method of randomisation: not stated. Concealment of allocation: not blinded for treatment allocation, blinded for outcome assessment Exclusions post-randomisation: 31 participants. Lost to follow-up: none.		
Participants	Country: not stated. Setting: hospital. No.: 197 participants. Age: mean 54 years. Sex: M:F 111:84. Inclusion criteria: informed consent. Exclusion criteria: two or more previous episodes of DVT or PE, currently active bleeding, active ulcers, known familial bleeding or coagulation disorder (i.e. known deficiency of antithrombin III, protein C or protein S), concurrent PE, treatment for the DVT with standard heparin lasting more than 48 hours, or impossibility of being or inability to be treated at home with LMWH or standard heparin. Also excluded were: people with neoplastic disorders requiring surgery or chemotherapy in the following 3 months, and those with likelihood of low or no compliance and/or inability to be included in a follow- up, pregnancy and a platelet count below 100,000 per mm ³		
Interventions	 Treatment: LMWH: administered primarily at home and body weight adjusted (nadroparin 0.1 mL per kg twice daily). Doses were 0.6, 0.8 and 1 mL (respectively equivalent to 6150, 8200 and 10,250 anti-factor Xa IU). Dose most suitable to the participant's weight was chosen Control: UFH: i.v. bolus of 5000 IU initially, followed by continuous infusion of 20, 000 IU. Dose was adjusted to maintain APTT between 60 and 85 seconds Treatment duration: LMWH: 5.1 days; UFH: 5.4 days. Oral anticoagulation: more than 3 months. 		
Outcomes	Primary: symptomatic or asymptomatic (detected by colour duplex scanning) recurrent DVT or DVT extension in 3 months after randomisation Secondary: bleeding during administration of the study medication or within 48 hours after discontinuation; PE; number of hospital days; number of participants treated di- rectly at home without hospital admission		
Notes	Follow-up: 3 months. 2 UFH groups (s.c. and i.v.).		
Risk of bias			
Bias	Authors' judgement Support for judgement		

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Belcaro 1999 (Continued)

Random sequence generation (selection bias)	Unclear risk	Insufficient information.
Allocation concealment (selection bias)	Unclear risk	Insufficient information.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	All reported outcome events were reviewed by a central panel unaware of the treatment assigned and participant's identity
Incomplete outcome data (attrition bias) All outcomes	Low risk	Missing outcome data balanced in numbers across intervention groups
Selective reporting (reporting bias)	Low risk	The published report includes all expected outcomes.
Other bias	Low risk	The study appears to be free of other sources of bias.

Breddin 2001

Methods	Study design: randomised controlled trial. Method of randomisation: stratified according to site. Concealment of allocation: not blinded for treatment allocation, blinded for outcome assessment Exclusions post-randomisation: none. Lost to follow-up: none.
Participants	Country: Argentina, Austria, Czech Republic, Denmark, Germany, Hungary, Israel, Poland, Norway, United Kingdom Setting: hospital. No.: 1137 participants. Age: mean 58 years. Sex: 621 males. Inclusion criteria: acute DVT confirmed by venography without symptoms lasting longer than 14 days Exclusion criteria: presence of thrombi only in isolated calf veins or isolated muscle veins; clinically symptomatic PE; treatment with UFH, LMWH, or VKA for 24 hours or more before enrolment; uncontrolled hypertension; stroke within 3 weeks of enrol- ment; cerebral vascular aneurysm or active gastroduodenal ulcer; bacterial endocarditis; thrombocytopenia (< 100,000 platelets/mm ³ ; severe liver or renal insufficiency; receipt of spinal or epidural anaesthesia or lumbar puncture in the 5 days before enrolment; surgery in the 5 days before enrolment; concomitant treatment with fibrinolytic agents

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Breddin 2001 (Continued)

	or platelet function inhibitors; a body weight of less than 35 kg; pregnancy and known drug abuse
Interventions	Treatment: LMWH: Reviparin (Clivarin, Knoll, Ludwigshafen, Germany) twice daily, body weight adjusted (7000 anti-factor Xa IU for a weight of 35 to 45 kg, 8400 IU for 46 to 60 kg and 12,600 IU for more than 60 kg) Control: 5000 IU i.v. UFH plus continuous i.v. infusion of 1250 IU/hour (dose-adjusted APTT × 1.5 to 2.5 Treatment duration: LMWH 5 to 7 days, UFH until INR > 2.0 (and maintained) Oral anticoagulation: in both groups (started day 1) for 90 days
Outcomes	Primary: change in venographically determined thrombus size (Marder's score) between base line and day 21 (± 2 days) Secondary: Clinical outcomes: recurrent DVT or PE during initial treatment and 3 months' follow- up; major haemorrhagic events between day 0 and 21
Notes	Follow-up: 90 days. LMWH once daily group (374 participants) not included in analysis because LMWH was given for 28 days

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Insufficient information.
Allocation concealment (selection bias)	Unclear risk	Insufficient information.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Blinded assessment of outcomes.
Incomplete outcome data (attrition bias) All outcomes	High risk	Reasons for missing second venogram and therefore exclusion for efficacy analysis are not provided and missing outcome data imbalanced in numbers across intervention groups
Selective reporting (reporting bias)	Low risk	The published report includes all expected outcomes.

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 30 thromboembolism (Review)

Breddin 2001 (Continued)

Other bias	Low risk	The study appears to be free of other sources of bias.		
Columbus 1997				
Methods	Method of randomisation: strat DVT only or with PE, accordi Concealment of allocation: no assessment	Exclusions post-randomisation: none.		
Participants	Setting: hospital. No.: 1021 participants. Age: mean 60. Sex: 525 males. Inclusion criteria: acute sympto DVT documented by ultrason lung scanning (high probability non-diagnostic, by demonstrati Exclusion criteria: therapeutic more than 24 hours; contraind therapy; gastrointestinal bleedin within the previous 3 days; a st	 No.: 1021 participants. Age: mean 60. Sex: 525 males. Inclusion criteria: acute symptomatic DVT and/or PE requiring antithrombotic therapy. DVT documented by ultrasonography or venography and PE by ventilation-perfusion lung scanning (high probability of PE), pulmonary angiography or, if lung scanning war non-diagnostic, by demonstrating DVT by compression ultrasonography or venograph Exclusion criteria: therapeutic doses of LMWH, UFH or oral anticoagulant therapy for more than 24 hours; contraindications for anticoagulant therapy; planned thrombolyti therapy; gastrointestinal bleeding in the preceding 14 days; surgery requiring anaesthesi within the previous 3 days; a stroke in the preceding 10 days; platelet count < 100,000 mm³; weight < 35 kg; pregnant or of childbearing potential and not using adequation. 		
Interventions	body weight adjusted fixed-dos left to treating physician Control: UFH: APTT-adjuste intravenous bolus of 5000 IU Treatment duration: at least 5 consecutive days	Control: UFH: APTT-adjusted dose, continuous i.v. infusion in hospital after initi intravenous bolus of 5000 IU Treatment duration: at least 5 days; treatment cessation if INR was 2.0 or above for consecutive days Oral anticoagulation: started on first or second day and continued for a total of 12 week		
Outcomes	randomisation Secondary: major haemorrhag	Primary: symptomatic DVT or PE during initial treatment and within 12 weeks of randomisation Secondary: major haemorrhage during initial treatment and within 12 weeks of ran- domisation; death within 12 weeks of randomisation		
Notes	PE: 138 (27%) versus 133 (26	Follow-up: 12 weeks. DVT only: LMWH 372 (73%) and UFH 378 (74%). PE: 138 (27%) versus 133 (26%). In retrospect, 3 participants with DVT only and 2 with PE should have been exclude		

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 31 thromboembolism (Review)

Columbus 1997 (Continued)

Risk of bias			
Bias	Authors' judgement	Support for judgement	
Random sequence generation (selection bias)	Low risk	Randomisation was performed with a com- puter algorithm.	
Allocation concealment (selection bias)	Low risk	Central allocation by a 24-hour telephone service.	
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes	
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Information on all suspected outcome events and deaths was reviewed and clas- sified by a central adjudication committee whose members were unaware of the treat- ment assignments	
Incomplete outcome data (attrition bias) All outcomes	Low risk	No missing outcome data.	
Selective reporting (reporting bias)	Low risk	The published report includes all expected outcomes.	
Other bias	Low risk	The study appears to be free of other sources of bias.	
Decousus 1998			
Methods	Study design: randomised controlled trial. Method of randomisation: stratified according to centre. Concealment of allocation: not blinded for treatment allocation, blinded for outcome assessment		

	Exclusions post-randomisation: none. Lost to follow-up: 4 (1 vital status; 3 for the assessment of non-fatal events)
Participants	Country: France (44 centres). Setting: hospital. No.: 400 participants. Age: mean 72. Sex: 190 males. Inclusion criteria: acute proximal DVT confirmed by venography with or without symptomatic PE; at high risk for PE

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Decousus 1998 (Continued)

	Exclusion criteria: placement of previous filter; contraindication to or failure of anticoag- ulant therapy; curative anticoagulant therapy lasting more than 48 hours; indication for thrombolysis; short life expectancy; allergy to iodine; hereditary thrombophilia; severe renal or hepatic failure; pregnancy; likelihood of non-compliance
Interventions	Treatment: LMWH: Enoxaparin (Rhone-Poulenc Rorer) body weight-adjusted fixed dose (1 mg per kg body weight), s.c., twice daily (100 anti-factor Xa IU per mg) Control: UFH: APTT-adjusted, continuous i.v. infusion (started with 500 IU per kg of body weight per day), after initial i.v. bolus dose of 5000 IU Treatment duration: 8 to 12 days; discontinuation if INR was 2 or more for 2 consecutive days Oral anticoagulation: warfarin or acenocoumarol started on day 4 and continued for at least 3 months
Outcomes	Primary: symptomatic or asymptomatic PE within the first 12 days after randomisation; all symptomatic recurrent VTE Secondary: major haemorrhage during the initial treatment period; mortality
Notes	Follow-up: 2 years. The outcome of recurrent VTE was only reported for a follow-up period of 3 months (also included as the incidence at the end of follow-up)

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Randomisation was performed by a com- puter system.
Allocation concealment (selection bias)	Low risk	Allocation was performed by a central 24- hour telephone system
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	All pulmonary investigations and all doc- umented symptomatic events, including deaths, were validated by an independent adjudication committee whose members were unaware of the treatment assignments
Incomplete outcome data (attrition bias) All outcomes	Low risk	Reasons for missing outcome data unlikely to be related to true outcome
Selective reporting (reporting bias)	Low risk	The published report includes all expected outcomes.

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Decousus 1998 (Continued)

Other bias	Unclear risk	The study has a potential source of bias due to the fact that 2 interventions (the effec- tiveness of a vena cava filter and the efficacy of LMWH) are investigated in the same population. There is insufficient Informa- tion about the number of participants with a vena cava filter across intervention groups
Faivre 1988		
Methods	Study design: randomised controlled trial. Method of randomisation: not stated. Concealment of allocation: not blinded for treatment allocation, blinded for outcome assessment Exclusion post-randomisation: 1 in UFH group (thrombocytopenia) Lost to follow-up: 9 participants had no second phlebography (3 CY 222, 6 UFH)	
Participants	Country: France. Setting: hospital. No.: 68 participants. Age: mean 66 years. Sex: 39 males. Inclusion criteria: symptomatic DVT and/or symptomatic PE, or symptomatic PE con- firmed by ventilation-perfusion scan and a positive phlebogram Exclusion criteria: > 2 weeks symptoms of DVT or PE with massive PE; extension of the thrombus into the inferior vena cava	
Interventions	IU and continued with body weight-adjust Xa IU/kg/24 hours), s.c., twice daily	
Outcomes	Primary: change in thrombus size (Marder Secondary: major haemorrhage during the	
Notes	LMWH and 34% of participants allocated Repeated venography; participants with th venographic evaluation.	nce of PE (66% of participants allocated to to UFH had a PE). rombotic and bleeding events excluded from riteria for diagnosis of recurrent VTE were

Risk of bias

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 34 thromboembolism (Review)

Faivre 1988 (Continued)

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Insufficient information.
Allocation concealment (selection bias)	Unclear risk	Insufficient information.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Blinded for outcome assessment.
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Insufficient information.
Selective reporting (reporting bias)	Unclear risk	Insufficient information.
Other bias	Unclear risk	Insufficient information. ? baseline differ- ences mentioned above?

Fiessinger 1996

Methods	Study design: randomised controlled trial. Method of randomisation: not stated. Concealment of allocation: not blinded for treatment allocation, blinded for outcome assessment Exclusions post-randomisation: 10 participants in dalteparin group and 5 participants in UFH group did not have DVT Lost to follow-up: 32 participants (13 versus 19) did not have a second phlebogram; 2 (1 versus 1) participants were considered not to have DVT; 20 participants (8 versus 12) were incorrectly included
Participants	Country: Austria, France, Spain and Sweden (16 centres). Setting: hospital. No.: 253 participants. Age: mean 61 years. Sex: 115 males. Inclusion criteria: distal and/or proximal DVT with 8 or more days of symptoms Exclusion criteria: clinical signs suggestive of PE; history of recent DVT (< 1 year) or sequelae of a previous DVT in the same leg; treatment with therapeutic doses of UFH or LMWH prior to randomisation; malignant hypertension; renal or hepatic insufficiency; platelet count < 100 x 10 ⁴ /litre; known hypersensitivity to contrast media; surgery within 5 days of starting treatment; intracerebral bleeding in previous 2 months, gastrointestinal

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Fiessinger 1996 (Continued)

	bleeding in previous 2 weeks; pregnancy/lactation	
Interventions	Treatment: LMWH: 1 mL active substance equivalent to 10,000 anti-factor Xa IU (Dalteparin, Fragmin) s.c. injection (200 IU/kg) o.d Bolus dose of 5000 IU. s.c. if randomisation before phlebography, otherwise a first full-dose Control: UFH: before phlebography: bolus dose of 5000 IU i.v. followed by continuous i.v. infusion of 20,000 to 40,000 IU/24 hours APTT-adjusted (1.5 to 3.0 ×). After phlebography a bolus i.v. injection administered prior to infusion of UFH at discretion of attending physician Treatment duration: 5 to 10 days, when the prothrombin time (INR) was within thera- peutic range (2 to 3) on 2 consecutive days Oral anticoagulation: started on day of inclusion or day after. Period determined by attending physician; mean period of treatment 5.3 months in both groups	
Outcomes	Primary: change in thrombus size (Marder's score); recurrent VTE during initial treat- ment (prospective follow-up) and at the end of 6 months' follow-up; PE during initial treatment and at the end of 6 months' follow-up Secondary: major haemorrhage during initial treatment; mortality; mortality in partici- pants with malignancy at entry	
Notes	20 participants not correctly included; 32 participants without second phlebography. Follow-up: 6 months, but 23 participants lost to follow-up; of these 13 were alive	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Insufficient information.
Allocation concealment (selection bias)	Unclear risk	Insufficient information.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Outcome assessment was blinded and the non-blinding of others is unlikely to intro- duce bias
Incomplete outcome data (attrition bias) All outcomes	High risk	Missing outcome data imbalanced in num- bers across intervention groups
Selective reporting (reporting bias)	Unclear risk	Insufficient information.

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 36 thromboembolism (Review)

Fiessinger 1996 (Continued)

Other bias	Low risk	The study appears to be free of other sources of bias.	
Findik 2002			
Methods	Study design: randomised contro Method of randomisation: not s Concealment of allocation: not l Exclusion post-randomisation: n Lost to follow-up: none.	tated. blinded for treatment allocation	
Participants	by ventilation-perfusion lung se indeterminate result accompanie Exclusion criteria: massive PE r traindication for anticoagulant t ticoagulant therapy at a therapeu	Setting: hospital. No.: 59 participants. Age: mean 50 years.	
Interventions	weight twice daily Control: UFH: Starting with a b v. infusion of 1000 IU/hour. UF Treatment duration: approximat	n s.c. 1 mg/kg, 100 anti-factor Xa IU per kg of body polus injection i.v. 5000 IU followed by a continuous i. H dose was adjusted (APTT-1.5 to 2.5 × control value) tely 7 days. the second day for a total of 6 months	
Outcomes	Primary: recurrent VTE, major h at 3 months	naemorrhage and mortality during initial treatment and	
Notes	Blinding for outcome assessmen	t was not reported.	
Risk of bias			

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Insufficient information.
Allocation concealment (selection bias)	Unclear risk	Insufficient information.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 37 thromboembolism (Review)

Findik 2002 (Continued)

		non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Insuffcient information.
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	No information about missing outcome data provided.
Selective reporting (reporting bias)	Low risk	Published report includes all expected out- comes.
Other bias	High risk	The low participant numbers in both LMWH and UFH arms and low event rates reduced the statistical power of the study to detect a significant difference between the arms
Goldhaber 1998		
Methods	Study design: randomised controlled trial. Method of randomisation: computerised, not stratified. Concealment of allocation: not blinded for treatment allocation, blinded for outcome assessment Exclusions post-randomisation: none. Lost to follow-up: not stated.	
Participants	Country: United States. Setting: hospital. No.: 81 participants. Age: mean 54 years. Sex: 43 males. Inclusion criteria: acute (within 14 days) symptomatic DVT of the legs documented by ultrasound and participants had to be deemed appropriate for discharge home Exclusion criteria: high-risk DVT involving 3 proximal veins; pelvic vein thrombosis; current symptomatic PE; expected prolonged hospitalisation for other reasons; haemo- globin < 85 g/litre or platelet count < 100 × 10° /litre	
Interventions	Treatment: LMWH: 130 anti-factor Xa IU/kg ardeparin sodium twice daily subcuta- neously for 5 to 15 days Control: UFH, heparin sodium 5000- to 7500-unit bolus followed by continuous i. v. administration to achieve APTT of 1.5 to 2.5. Titration guided by Cruickshank nomogram Treatment duration: LMWH 5 to 15 days, UFH 5 days or more to achieve target APTT Oral anticoagulation: 6 weeks.	

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 38 thromboembolism (Review)

Goldhaber 1998 (Continued)

Outcomes	Primary: change in thrombus size; recurrent DVT or PE. Secondary: major and minor haemorrhage.	
Notes	Repeated venography at the end of follow-up (6 weeks).	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Randomisation and allocation was accom- plished by calling a central computerised service
Allocation concealment (selection bias)	Low risk	Randomisation and allocation was accom- plished by calling a central computerised service
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Insufficient information.
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	No reasons for missing data provided.
Selective reporting (reporting bias)	Low risk	Published report includes all expected out- comes.
Other bias	Low risk	The study appears to be free of other sources of bias.

Harenberg 2000a

Methods	Study design: randomised controlled trial. Method of randomisation: not stated. Concealment of allocation: blinded for outcome assessment. Exclusions post-randomisation: not stated. Lost to follow-up: not stated.
Participants	Country: Austria, Germany, Switzerland, Czech Republic. Setting: hospital. No.: 541 were eligible of which 3 withdrew informed consent; therefore 538 participants were assigned

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 39 thromboembolism (Review)

Harenberg 2000a (Continued)

	Age: 30 years and older. Sex: Males and females (breakdown not supplied). Inclusion criteria: acute symptomatic proximal DVT (thrombosis of the popliteal vein or proximal) documented by ascending venography Exclusion criteria: indication for surgical or fibrinolytic treatment of DVT; duration of symptoms for more than 3 weeks; ongoing oral anticoagulation; renal failure; severe hy- pertension (> 200 mmHg systolic and > 105 mmHg diastolic while on antihypertensive treatment); severe hepatic failure; currently active bleeding or disorders contraindicating anticoagulant therapy; contraindication to oral anticoagulants; pregnancy; known intol- erance to heparins; intolerance to contrast media; any operation within the past 8 days; acute severe PE; platelet count < 100,000/µL; treatment with heparin > 24 hours before inclusion; treatment with platelet-inhibiting drugs (100 mg or more acetylsalicylic acid daily allowed)
Interventions	Treatment: LMWH: fixed dose 8000 anti-factor Xa IU (Certoparin) s.c., twice daily Control: UFH: adjusted to APTT 2 to 3 × the reference value. Treatment duration: 7 to 15 days. Oral anticoagulation: at least 6 months.
Outcomes	Primary: change in thrombus size (Marder's score), recurrent VTE, major bleeding and death during treatment and after 6 months' test follow-up
Nata	

- Notes
- Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Insufficient information.
Allocation concealment (selection bias)	Unclear risk	Insufficient information.
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Insufficient information.
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Data on all potential outcome events were evaluated by an independent committee, which was unaware of the treatment assign- ment
Incomplete outcome data (attrition bias) All outcomes	Low risk	Missing outcome data balanced in numbers across intervention groups
Selective reporting (reporting bias)	Low risk	Published report includes all expected out- comes.

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 40 thromboembolism (Review)

Harenberg 2000a (Continued)

Other bias	High risk	The study was sponsored by Novartis Phar- macological GmbH, Nuremberg, Ger- many	
Hull 1992			
Methods	Method of randomisation: co centre Concealment of allocation: bli	Concealment of allocation: blinded for treatment allocation and outcome assessment Exclusions post-randomisation: none.	
Participants	Setting: hospital. No.: 432 participants. Age: 161 participants under 6 Sex: 140 males, 291 females. Inclusion criteria: symptomati tomatic PE Exclusion criteria: active bleec allergy to heparin, bisulphites episodes of DVT or PE; histo thrombocytopenia; severe mal systolic and 130 mmHg or mo thy); severe renal failure; requi dance at follow-up visits. Eligi ment with warfarin, LMWH o	No.: 432 participants. Age: 161 participants under 60 years, 270 participants over 60 years Sex: 140 males, 291 females. Inclusion criteria: symptomatic or asymptomatic proximal DVT with or without symp-	
Interventions	kg, s.c., o.d Control: UFH: dose-adjusted 40,320 Units/24 hours; or in p of 5000 Units Treatment duration: 6 days pr Oral anticoagulation: warfarin	Control: UFH: dose-adjusted APTT × 1.5 to 2.5, continuous i.v. infusion starting with 40,320 Units/24 hours; or in people at high risk, 29,760 Units/24 hours. Initial i.v. bolus	
Outcomes	treatment	Primary: recurrent DVT and PE; major haemorrhage during or immediately after initial treatment Secondary: minor haemorrhage; mortality.	
Notes	Placebo controlled. Follow-up: 3 months. More women in UFH group;		

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 41 thromboembolism (Review)

Hull 1992 (Continued)

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	A randomised computer-derived treatment schedule was used to assign the participants to the treatment group
Allocation concealment (selection bias)	Low risk	A randomised computer-derived treatment schedule was used to assign the participants to the treatment group
Blinding of participants and personnel (performance bias) All outcomes	Low risk	A double-blind clinical trial. Participants received either intravenous UFH with subcutaneous placebo or subcutaneous LMWH with intravenous placebo
Blinding of outcome assessment (detection bias) All outcomes	Low risk	A double-blind clinical trial. The outcome assessment was blinded
Incomplete outcome data (attrition bias) All outcomes	Low risk	No missing outcome data.
Selective reporting (reporting bias)	Low risk	Published report includes all expected out- comes.
Other bias	Low risk	There were more women in the intravenous heparin group. To assess the possible effec of this potential gender imbalance, multi- ple logistic regression was used. No signif- icant effect was found. The study seems to be free of other sources of bias

Kakkar 2003

Methods	Study design: multicentre, randomised, open-label, parallel group comparison trial Method of randomisation: not stated Concealment of allocation: not stated Exclusions post-randomisation: 54 participants Lost to follow-up: none
Participants	Country: Spain, Poland and United Kingdom Setting: 27 hospitals No.: 324 participants: 94 bemiparin, 105 bemiparin + VKA, 98 UFH Age: bemiparin mean 63.2 (45.1 to 70.8) years, bemiparin + VKA mean 61.2 (44.4 to 69.5) years, UFH mean 61.2 (49.9 to 70.5) years, Sex: bemiparin 58 M/36 F, bemiparin + VKA 61 M/44 F, UFH 63 M/35 F,

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 42 thromboembolism (Review)

Kakkar 2003 (Continued)

Inclusion criteria: people with an acute DVT of the legs, confirmed by venography and who had symptoms for no more than 14 days Exclusion criteria: people reviewing therapeutic doses of heparin or a vitamin K antagonist for more than 48 hours prior to enrolment, clinically symptomatic pulmonary embolism, pregnancy confirmed by urine analysis, ischarenic cerebral vascular accident 1 month prior to enrolment, known cerebral vascular anceursm, active duodenal ulcer or bacterial endocarditis, severe liver or renal failure, spinal or epidural anaesthesia or lumbar puncture 3 days prior to enrolment, uncontrolled hypertension, allergy to heparin, warfarin, sodium or iodinated contrast medium, history of heparin-associated thrombocytopenia or platelet count of less than 100,000 platelets per mm³, concurrent treatment with fibrinolytic agents, a body weight of less than 35 kg, treatment with an investigational drug in the last 4 weeks prior to enrolment, inability to attend follow-up due to geographic inaccessibility and known drug use Interventions Treatment 1: 115 anti-Xa IU per kg of bemiparin as 1 injection every 24 hours based on participants' weight (5000 anti-Xa for weight < 50 kg, 7500 anti-Xa for weight 50 to 70 kg and 10,000 anti-Xa 10 for more than 70 kg) followed by VKA from day 310 mg per day for first 3 days then adjusted to achieve an INR between 2 and 3 for 12 weeks Iteratment 2: 115 anti-Xa IU per kg of bemiparin as 1 injection every 24 hours based on participants' weight 5000 anti-Xa for weight < 50 kg, 7500 anti-Xa for weight 50 to 70 kg and 10,000 anti-Xa 10 for more than 70 kg) followed by VKA from day 310 mg per day for first 3 days then adjusted to achieve an 100 kg torologinat a dose of 40, 000 TU per 24 hours in participants at low risk of bleeding and 30,000 TU per 24 hours in participants at low risk of bleeding and 30,000 TU per 24 hours in parti		
participants' weight (5000 anti-Xa for weight < 50 kg, 7,500 anti-Xa for weight 50 to 70 kg and 10,000 anti-Xa IU for more than 70 kg) followed by VKA from day 3 10 mg per day for first 3 days then adjusted to achieve an INR between 2 and 3 for 12 weeks Treatment 2: 115 anti-Xa IU per kg of bemiparin as 1 injection every 24 hours based on participants' weight (5000 anti-Xa for weight < 50 kg, 7500 anti-Xa for weight 50 to 70 kg and 10,000 anti-Xa IU for more than 70 kg) followed by fixed daily dose of 3500 anti-Xa units for 90 days Control: i.v. bolus of 5000 UFH followed by a continuous i.v. infusion at a dose of 40, 000 IU per 24 hours in participants at low risk of bleeding and 30,000 IU per 24 hours in participants at high risk of bleeding followed by VKA from day 3 10 mg per day for first 3 days then adjusted to achieve an INR between 2 and 3 for 12 weeks Treatment duration: 12 weeks.OutcomesPrimary: venographically confirmed change in thrombus size between baseline and day 14 assessed with the use of the Marder score and patency of deep venous system determined by venography or Doppler ultrasound at 12 weeks Secondary: symptomatic recurrence of DVT and PE, major bleeding (clinically overt and associated with a fall in haemoglobin level of at least 2.0 g per decilitre) and deathNotesFollow-up: 7 days, 14 days, 12 weeks and 28 weeks. In this 3-armed trial, 2 bemiparin groups (treatment 2), participants did not receive con- comitant VKA therapy. All other studies included in this group of participants VKA therapy and in order for our results to be comparable, data for this group of participants		who had symptoms for no more than 14 days Exclusion criteria: people receiving therapeutic doses of heparin or a vitamin K antag- onist for more than 48 hours prior to enrolment, clinically symptomatic pulmonary embolism, pregnancy confirmed by urine analysis, ischaemic cerebral vascular accident 1 month prior to enrolment, known cerebral vascular aneurysm, active duodenal ul- cer or bacterial endocarditis, severe liver or renal failure, spinal or epidural anaesthesia or lumbar puncture 3 days prior to enrolment, uncontrolled hypertension, allergy to heparin, warfarin, sodium or iodinated contrast medium, history of heparin-associated thrombocytopenia or platelet count of less than 100,000 platelets per mm ³ , concurrent treatment with fibrinolytic agents, a body weight of less than 35 kg, treatment with an investigational drug in the last 4 weeks prior to enrolment, inability to attend follow-up
assessed with the use of the Marder score and patency of deep venous system determined by venography or Doppler ultrasound at 12 weeks Secondary: symptomatic recurrence of DVT and PE, major bleeding (clinically overt and associated with a fall in haemoglobin level of at least 2.0 g per decilitre) and deathNotesFollow-up: 7 days, 14 days, 12 weeks and 28 weeks. In this 3-armed trial, 2 bemiparin groups were compared with an UFH control group. However, in 1 of the bemiparin groups (treatment 2), participants did not receive con- comitant VKA therapy. All other studies included in this group of participants	Interventions	participants' weight (5000 anti-Xa for weight < 50 kg, 7,500 anti-Xa for weight 50 to 70 kg and 10,000 anti-Xa IU for more than 70 kg) followed by VKA from day 3 10 mg per day for first 3 days then adjusted to achieve an INR between 2 and 3 for 12 weeks Treatment 2: 115 anti-Xa IU per kg of bemiparin as 1 injection every 24 hours based on participants' weight (5000 anti-Xa for weight < 50 kg, 7500 anti-Xa for weight 50 to 70 kg and 10,000 anti-Xa IU for more than 70 kg) followed by fixed daily dose of 3500 anti-Xa units for 90 days Control: i.v. bolus of 5000 UFH followed by a continuous i.v. infusion at a dose of 40, 000 IU per 24 hours in participants at low risk of bleeding and 30,000 IU per 24 hours in participants at low risk of bleeding and 30,000 IU per 24 hours in participants at high risk of bleeding followed by VKA from day 3 10 mg per day for first 3 days then adjusted to achieve an INR between 2 and 3 for 12 weeks
In this 3-armed trial, 2 bemiparin groups were compared with an UFH control group. However, in 1 of the bemiparin groups (treatment 2), participants did not receive con- comitant VKA therapy. All other studies included in this review used concomitant VKA therapy and in order for our results to be comparable, data for this group of participants	Outcomes	assessed with the use of the Marder score and patency of deep venous system determined by venography or Doppler ultrasound at 12 weeks Secondary: symptomatic recurrence of DVT and PE, major bleeding (clinically overt
	Notes	In this 3-armed trial, 2 bemiparin groups were compared with an UFH control group. However, in 1 of the bemiparin groups (treatment 2), participants did not receive con- comitant VKA therapy. All other studies included in this review used concomitant VKA therapy and in order for our results to be comparable, data for this group of participants

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Quote: "Patients were randomly assigned". Comment: insufficient information to per- mit judgement.

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 43 thromboembolism (Review)

Kakkar 2003 (Continued)

Allocation concealment (selection bias)	Unclear risk	Quote: "Patients were randomly assigned". Comment: insufficient information to per- mit judgement.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Open label Quote: "The venograms were indepen- dently assessed by two radiologists of an in- dependent committee who were unaware of the patients treatment assignments" Comment: outcome assessors were blinded.
Incomplete outcome data (attrition bias) All outcomes	High risk	324 participants in intention-to-treat group but only 297 participants included in the per protocol population and only 255 followed up to day 84. Numbers lost to follow-up not adequately reported
Selective reporting (reporting bias)	Low risk	Published report includes all expected out- comes.
Other bias	High risk	The study was sponsored by Laboratorios Farmaceuticos Rovi, Madrid, Spain

Kirchmaier 1998

Methods	Study design: randomised controlled trial. Method of randomisation: computerised. Concealment of allocation: partly blinded for treatment allocation Exclusions post-randomisation: 6 participants. Lost to follow-up: none.	
Participants	Lost to ronow-up: none. Country: Austria, Czech Republic, Germany (total 23 centres) Setting: hospital. No.: 257 participants. Age: median 61 years. Sex: 133 males. Inclusion criteria: symptomatic DVT of the lower leg. Exclusion criteria: thrombi only in 1 or 2 calf veins; treatment with vitamin K antagonists; use of contrast media; surgery in the previous week; thrombocytopenic (< 100,000/µL)	

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 44 thromboembolism (Review)

Kirchmaier 1998 (Continued)

Interventions	Treatment: subcutaneous LMWH (certoparin) 8000 IU/kg twice daily Control: UFH: initial bolus of 5000 IU followed by 20 IU/kg/hour In both groups phenprocoumon was started between day 12 and 14. Heparin was stopped until an INR range between 2.0 and 3.5 was reached Treatment duration: at least 14 days. Oral anticoagulation: Oral anticoagulant therapy was continued for at least 6 months
Outcomes	Primary: recurrent VTE; major haemorrhage during initial treatment; change in throm- bus size; mortality at the end of follow-up
Notes	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Randomisation performed by a statistician, but there is insufficient information about the sequence generation
Allocation concealment (selection bias)	Low risk	Allocation was performed centrally by tele- phone.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Partly blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Outcome assessment by an investigator, who was blinded to the treatment the par- ticipants had received
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Reasons for missing phlebograms and per- fusion scans were not provided
Selective reporting (reporting bias)	Low risk	Published report includes all expected out- comes.
Other bias	Low risk	The study appears to be free of other sources of bias.

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 45 thromboembolism (Review)

Koopman 1996

Methods	Study design: randomised controlled trial. Method of randomisation: stratified according to centre. Concealment of allocation: not blinded for treatment allocation, blinded for outcome assessment Exclusions post-randomisation: none. Lost to follow-up: none.
Participants	Country: Netherlands, France, Italy, Australia, New Zealand. Setting: hospital. No.: 400 outpatients. Age: Mean 61 years. Sex: 203 males. Inclusion criteria: acute symptomatic proximal DVT documented by venography and/ or ultrasonography Exclusion criteria: VTE in last 2 years; suspected PE; previous treatment with heparin > 24 hours; life expectancy < 6 months; post-thrombotic syndrome; geographic inaccessibility
Interventions	Treatment: LMWH (Nadroparin-Ca, Fraxiparine) in body weight-adjusted fixed dose, s.c., twice daily. If appropriate, at home Control: UFH: APTT-adjusted dose, continuous i.v. infusion in hospital after initial i. v. bolus of 5000 Units Treatment duration: at least 5 days; treatment cessation if INR was 2.0 or above in 2 measurements 24 hours apart Oral anticoagulation: started on first day and continued for 3 months unless persistence of risk factors required its continuation beyond that period. INR 2.0 to 3.0
Outcomes	Primary: symptomatic recurrent VTE (DVT or PE) during initial treatment, after 3 months' follow-up and at the end of follow-up (6 months); major haemorrhage during initial treatment and after 3 months of follow-up Secondary: minor haemorrhage or death during initial treatment, after 3 months of follow-up and at the end of follow-up (6 months); other potential outcome events; quality of life
Notes	Follow-up: 6 months.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Randomisation and allocation was achieved by means of a central 24-hour telephone service
Allocation concealment (selection bias)	Low risk	Randomisation and allocation was achieved by means of a central 24-hour telephone service

Koopman 1996 (Continued)

Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Documentation of all potential outcome events was submitted to an independent adjudication committee whose members were unaware of the treatment assignments
Incomplete outcome data (attrition bias) All outcomes	Low risk	Missing outcome data balanced in numbers across intervention groups
Selective reporting (reporting bias)	Low risk	Published report includes all expected out- comes.
Other bias	Low risk	Study appears to be free of other sources of bias.

Leizorovicz 2011

Methods	Study design: international, multicentre, centrally randomised, open, parallel-group study Method of randomisation: computer generated randomisation scheme in a 1:1 ratio with central telephone randomisation Concealment of allocation: no allocation concealment mechanism was attempted Exclusions post-randomisation: none Lost to follow-up: none
Participants	Country: 8 countries (Belgium, France, Germany, Spain, Serbia, Croatia, Romania and Poland) Setting: 109 hospitals No.: 269 tinzaparin, 270 UFH Age: tinzaparin mean 82.9 \pm 5.7 years, UFH mean 82.6 \pm 5.8 years Sex: tinzaparin 92 M/177 F, UFH 102 M/168 F Inclusion criteria: people \geq 70 years with an acute objectively confirmed (compression ultrasonography or venography) symptomatic proximal or distal lower limb DVT or asymptomatic DVT if proximal and associated with a PE Exclusion criteria: people who had received treatment doses of heparins or thrombolytic agents within the previous 4 weeks prior to randomisation, received oral anticoagulation within the preceding week, planned use of high doses of acetylsalicylic acid (> 300mg/ day) or an NSAID, had a requirement for thrombolytic therapy, end stage renal dis- ease requiring dialysis, hepatic insufficiency, bacterial endocarditis, planned epidural or spinal anaesthesia, planned or recent (within 2 weeks) surgery, thrombocytopenia, severe uncontrolled hypertension, overt bleeding or recent stroke

Leizorovicz 2011 (Continued)

Interventions	Treatment: tinzaparin 175 IU/kg subcutaneous injection once daily Control: UFH (50 IU/kg i.v. bolus followed by twice daily subcutaneous injections in initial doses between 400 to 600 IU/kg/day then adjusted by APTT) Treatment duration: 5 days. Oral anticoagulation: VKA treatment initiated between days 1 and 3 and continued until at least day 90 ± 5
Outcomes	Primary: clinically relevant bleeding by day 90 \pm 5. Secondary: symptomatic recurrent VTE prior to day 90 \pm 5, major and minor bleedings prior to day 90 \pm 5, heparin-induced thrombocytopenia and death
Notes	Follow-up: 90 ± 5 days. Study was unexpectedly terminated early as at a predefined interim analysis conducted after completion of 350 participants, an excess mortality was observed in the tinzaparin group

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "Treatment assignment was pre- planned according to a computer generated randomisation scheme in a 1:1 ratio with central telephone randomisation" Comment: low risk of bias.
Allocation concealment (selection bias)	Unclear risk	Quote: "No allocation concealment mech- anism was attempted as the study was open" Comment: insufficient information to per- mit judgement.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Quote: "The study was open but care was taken to ensure that outcome assessors and data analysts were kept blinded to the allo- cation" Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Outcome assessors and data analysts were kept blinded to the allocation
Incomplete outcome data (attrition bias) All outcomes	Low risk	No missing outcome data.

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 48 thromboembolism (Review)

Leizorovicz 2011 (Continued)

Selective reporting (reporting bias)	Low risk	Published report includes all expected out- comes.	
Other bias	Unclear risk	Study sponsored by LEO Pharma.	
Levine 1996			
Methods	Method of randomisation: stra or ultrasonography), and cates at night, hospitalised) Concealment of allocation: no assessment	Concealment of allocation: not blinded for treatment allocation, blinded for outcome assessment Exclusions post-randomisation: none.	
Participants	peptic ulcer disease; familial b heparin treatment; inability to ing condition (e.g. cancer, inf to make follow-up visits becau	down not supplied).	
Interventions	Treatment: LMWH: enoxaparin (Rhone-Poulenc Rorer) body weight-adjusted fixed dose (1 mg/kg body weight), s.c., twice daily, at home. 1 vial: 1 mL/100 mg = 100 anti-factor Xa IU/mg) Control: UFH: APTT-adjusted, continuous i.v. infusion (started with 20,000 Units in 500 mL of 5% dextrose solution) in hospital after an initial i.v. bolus of 5000 Units Treatment duration: at least 5 days; discontinuation if INR was 2 or above and maintained for 2 consecutive days Oral anticoagulation warfarin sodium started on day 2 and continued for 3 months		
Outcomes	Primary: symptomatic recurrent VTE within 90 days of follow-up; major haemorrhage during the initial treatment or 48 hours after treatment cessation Secondary: minor haemorrhage; mortality.		
Notes	Some participants received 1 or 2 days UFH before randomisation; this was considered part of the overall duration of heparin treatment. Follow-up: 3 months.		

Risk of bias

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 49 thromboembolism (Review)

Levine 1996 (Continued)

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Insufficient information.
Allocation concealment (selection bias)	Low risk	Allocation over the telephone from a cen- tral site.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	All reported outcome events were reviewed by a central adjudication committee whose members were unaware of the treatment assignments
Incomplete outcome data (attrition bias) All outcomes	Low risk	No missing outcome data.
Selective reporting (reporting bias)	Low risk	Published report includes all expected out- comes.
Other bias	Low risk	The study appears to be free of other sources of bias.

Lindmarker 1994

Methods	Study design: randomised controlled trial. Method of randomisation: centrally organised using sealed envelopes and stratified for centre Concealment of allocation: not blinded for treatment allocation, blinded for outcome assessment Exclusions post-randomisation: 6 (1 UFH versus 5 fragmin). Lost to follow-up: for venographic assessment 18 (13 UFH versus 5 fragmin); for clinical outcome assessment 16 participants
Participants	Country: Sweden. Setting: hospital. No.: 204 outpatients. Age: mean 61 years. Sex: 116 males. Inclusion criteria: symptomatic distal and proximal DVT. Exclusion criteria: UFH treatment already given for more than 24 hours; surgery < 5 days before; previous DVT in the ipsilateral leg; suspected or verified PE; thrombectomy or thrombolysis indicated; DVT proximal of inguinal arch; intracranial bleeding within

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 50 thromboembolism (Review)

Lindmarker 1994 (Continued)

	previous 2 weeks; known haemorrhagic diathesis or disorders; platelet count below 100 \times 10° /litre; renal insufficiency (S-creatinine < 300 μ M); hepatic insufficiency with a prothrombin time < 40% (INR > 1.5); allergy to UFH, fragmin or contrast media; pregnancy or breastfeeding; severe hypertension
Interventions	Treatment: initial i.v. bolus injection of UFH 5000 Units followed by continuous i.v. infusion of UFH 800 to 1700/hour for a maximum of 24 hours after randomisation: LMWH (fragmin) body weight-adjusted fixed dose of 200 anti-factor Xa IU/kg with a maximum of 18,000 IU, s.c., o.d Control: initial i.v. bolus injection of UFH 5000 Units followed by continuous i.v. infusion of UFH 800 to 1700/hour; after randomisation: continuation of i.v. infusion with UFH dose-adjusted APTT × 1.5 to 3.0 Treatment duration: at least 5 days; treatment cessation if INR was within therapeutic range (2.0 to 3.0) for 2 consecutive days. Treatment duration no longer than 9 days Oral anticoagulation: warfarin sodium started on the day that venography was carried out and continued for a minimum of 3 months; INR 2.0 to 3.0
Outcomes	Primary: change in thrombus size (Marder's score); recurrent VTE; major haemorrhage Secondary: mortality; mortality in participants with malignant disease
Notes	Repeated venography on day 1 and within 4 days after discontinuation of heparin therapy. Follow-up: 6 months.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Insufficient information.
Allocation concealment (selection bias)	Unclear risk	Insufficient information.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Outcome assessment was blinded.
Incomplete outcome data (attrition bias) All outcomes	High risk	The participants who died or had a recur- rent VTE were not included in the analyses which may result in an underestimation of the number of participants with extended or unchanged thrombosis

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 51 thromboembolism (Review)

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Lindmarker 1994 (Continued)

Selective reporting (reporting bias)	High risk	Participants who died or had a recurrent VTE were not included in the analyses
Other bias	Low risk	The study appears to be free of other sources of bias.
Lopaciuk 1992		
Methods	Study design: randomised controlled trial. Method of randomisation: sealed envelopes, stratified for site of DVT Concealment of allocation: not blinded for treatment allocation, blinded for outcome assessment Exclusions post-randomisation: 3 participants in UFH group judged to be ineligible (2 with recent history of DVT and 1 deficient in antithrombin III) Lost to follow-up: 6 in LMWH group and 6 in UFH group (poor phlebogram, 6; absent phlebogram, 4; protocol violation (treatment for 15 days), 1; major bleeding with treatment cessation, 1)	
Participants	Country: Poland (6 centres). Setting: hospital. No.: 149 participants of which 117 participants had proximal DVT Age: mean 48 years. Sex: 81 males. Inclusion criteria: symptomatic proximal or calf DVT (phlebographically proven) Exclusion criteria: clinically suspected PE; phlegmasia caerulea dolens; treatment with heparin or oral anticoagulants prior to admission; history of VTE in previous 2 years; surgery or trauma within previous 3 days; contraindication to heparin therapy; preg- nancy; documented antithrombin III deficiency	
Interventions	Treatment: LMWH: fraxiparine fixed dose: 92 anti-factor Xa IU/kg, s.c., twice daily Control: UFH: initial i.v. bolus of 5000 IU followed by 250 IU/kg s.c., twice daily; dose- adjusted APTT × 1.5 to 2.5 s.c Treatment duration: 10 days. Oral anticoagulation: acenocoumarol started on day 7 and continued for at least 3 months; INR 2.0 to 3.0	
Outcomes	Primary: change in thrombus size (Arnesen score); recurrent DVT; PE Secondary: major and minor haemorrhage; mortality; mortality in participants with malignant disease	
Notes	Proximal DVT: 58 (LMWH) versus 59 (UFH). Distal DVT: 16 (LMWH) versus 13 (UFH). 12 participants excluded from repeated venography analysis. Follow-up: 3 months.	
Risk of bias		
Bias	Authors' judgement	Support for judgement

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 52 thromboembolism (Review)

Lopaciuk 1992 (Continued)

Random sequence generation (selection bias)	Unclear risk	Insufficient information.
Allocation concealment (selection bias)	Low risk	Sealed envelopes were used.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Blind evaluation of phlebographic results.
Incomplete outcome data (attrition bias) All outcomes	Low risk	Missing outcome data balanced in numbers across intervention groups
Selective reporting (reporting bias)	Unclear risk	Insufficient information.
Other bias	High risk	There was an imbalanced exclusion at base- line.

Luomanmaki 1996

Methods	Study design: randomised controlled trial. Method of randomisation: not stated. Concealment of allocation: not blinded for treatment allocation and for clinical outcome assessment; blinded for assessment of venograms at entry to study and at the end of the initial treatment period Exclusions post-randomisation: 78 randomised participants excluded because DVT found not to be present after randomisation Lost to follow-up: no information given.
Participants	Country: Sweden and USA (2 centres). Setting: hospital. No.: 248 participants. Age: mean 57.5 years (LMWH); mean 60.5 years (UFH). Sex: 125 males. Inclusion criteria: clinically suspected or verified DVT. Exclusion criteria: none stated.
Interventions	Treatment: LMWH: dalteparin fixed dose body weight-adjusted (200 IU/kg), s.c., o.d Control: UFH: dose-adjusted APTT × 1.5 to 3.0, continuous i.v. infusion Treatment duration: 5 to 10 days until therapeutic effect of oral anticoagulants was reached Oral anticoagulation: started during the initial heparin treatment

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 53 thromboembolism (Review)

Luomanmaki 1996 (Continued)

Outcomes	Primary: change in thrombus size (Marder's score); recurrent VTE (no blind assessmen	
	; major haemorrhage; mortality at the end of follow-up	

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Randomisation was conducted using a Sta- tistical Analysis System Program
Allocation concealment (selection bias)	High risk	No central allocation: randomisation was conducted separately at each participating centre
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Blinded evaluations of venograms.
Incomplete outcome data (attrition bias) All outcomes	Low risk	Reasons for missing outcome data unlikely to be related to true outcome
Selective reporting (reporting bias)	Low risk	Published report includes all expected out- comes.
Other bias	High risk	There was a significantly higher incidence of malignancy in participants randomised to UFH

Merli 2001

Methods	Study design: randomised controlled trial. Method of randomisation: block randomisation without stratification Concealment of allocation: partly blinded for treatment allocation, blinded for outcome assessment Exclusions post-randomisation: not stated. Lost to follow-up: not stated.
Participants	Country: Australia, Austria, Belgium, Denmark, France, Hungary, Ireland, Israel, Italy, Netherlands, Norway, Poland, Spain, Sweden, United Kingdom and USA Setting: hospital.

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 54 thromboembolism (Review)

Merli 2001 (Continued)

	No.: 900 participants. Age: mean 61 years. Sex: 492 males. Inclusion criteria: symptomatic lower extremity DVT confirmed by venography or ul- trasonography (if venography was inconclusive), symptomatic PE confirmed by high probability ventilation-perfusion scanning or positive pulmonary angiography with con- firmation of lower extremity DVT. All those who were eligible underwent baseline lung scanning or angiography Exclusion criteria: more than 24 hours of previous treatment with heparin or warfarin; need for thrombolytic therapy; known haemorrhagic risk, including active haemorrhage, active intestinal ulcerative disease, known angiodysplasia or eye, spine or central ner- vous system surgery within the previous month; renal insufficiency (serum creatinine concentration > 180 µmol/litre (2.03 mg/dL)); severe hepatic insufficiency; allergy to heparin, protamine, porcine products, iodine or contrast media; history of heparin-as- sociated thrombocytopenia or heparin- or warfarin-associated skin necrosis; treatment with other investigational therapeutic agents within the previous 4 weeks; inferior vena cava interruption; known pregnancy or lactation
Interventions	Treatment: LMWH: enoxaparin weight-adjusted s.c. dose (1.0 mg/kg of body weight twice daily or 1.5 mg/kg of body weight o.d.) Control: UFH: initial i.v. bolus injection followed by an infusion based on an approved nomogram. In general: 6 hours after initial bolus an adjusted dose was given to maintain APTT between 55 and 80 seconds. APTT was measured daily Treatment duration: enoxaparin and heparin treatment were continued for at least 5 days, and warfarin was started within 72 hours of initial study drug administration. 43 participants received phenprocoumon in place of warfarin sodium. INR between 2.0 and 3.0 Oral anticoagulation: oral anticoagulation was continued for at least 3 months
Outcomes	Primary: worsening or recurrence of DVT or PE within 3 months Secondary: clinical overt minor or major haemorrhage.
Notes	Participants who received LMWH (2 groups; o.d. and twice daily) were analysed as 1 group

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	The authors refer to a random number table.
Allocation concealment (selection bias)	Low risk	The randomisation numbers were affixed to sealed treatment kits
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Partly blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 55 thromboembolism (Review)

Merli 2001 (Continued)

		was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	The Outcome Adjudication Committee provided blinded outcome assessments
Incomplete outcome data (attrition bias) All outcomes	Low risk	Missing outcome data balanced in num- bers across intervention groups, with simi- lar reasons for missing data across groups
Selective reporting (reporting bias)	Low risk	Published report includes all expected out- comes.
Other bias	Low risk	The study appears to be free of other sources of bias.

Meyer 1995

Methods	Study design: randomised, multicentre pilot study. Method of randomisation: not stated. Concealment of allocation: sealed envelopes, not blinded for treatment allocation, blinded for outcome assessment Exclusions post-randomisation: none. Lost to follow-up: none.
Participants	Country: France. Setting: hospital. No.: 60 participants: 29 LMWH, 31 UF heparin. Age: mean 60 (range 26 to 84) years LMWH, mean 61 (20 to 88) years UF heparin Sex: LMWH 9 M/20 F, UF heparin 17 M/14 F Inclusion criteria: men and women > 18 years, weighing 45 to 90 kg and with onset of symptoms suggestive of acute PE within the 5 preceding days Exclusion criteria: known pregnancy or breastfeeding, major surgical procedure or organ biopsy within the last 5 days, ischaemic cerebrovascular accident within the past 30 days or cerebral haemorrhage within the last 3 months, known haemorrhagic diathesis, active peptic ulcer, pre-existing significant cardiorespiratory disease, known proliferative diabetic retinopathy, known allergy to heparin or contrast media, platelet count < 100 10 ⁶ /L, chronic renal failure, chronic liver disease, treatment with UFH or LMWH at full dosage for more than 24 hours before randomisation, planned hospital stay < 10 days, oral anticoagulant therapy within 5 days before randomisation and any clinical condition which in the opinion of the physician in charge would not allow safe fulfilment of the protocol
Interventions	Treatment: LMWH: fragmin at a fixed dose of 120 anti-Xa IU/kg subcutaneously twice daily and without any laboratory adjustment Control: UFH as a continuous intravenous infusion at an initial dosage of 500 IU/kg/ 24 hours and adjusted daily to maintain APTT between 2 to 3 times the control value Treatment duration: 10 days Oral anticoagulation: acenocoumarol started on day 7 and continued for at least 3 months

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 56 thromboembolism (Review)

Meyer 1995 (Continued)

Outcomes	Primary: incidence of PE recurrence within the first 10 days of treatment Secondary: pulmonary scintigraphic vascular obstruction score (PVOS), major bleeding	
Notes	Follow-up: 3 months	
Risk of bias		
Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Quote: "Treatment was randomly allo- cated". Comment: insufficient information to per- mit judgement.
Allocation concealment (selection bias)	Unclear risk	Quote: "Treatment was randomly allocated using sealed envelopes" Comment: although the use of assignment envelopes is described, it remains unclear whether envelopes were sequentially num- bered and opaque
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "Open study. All angiograms were reviewed and scored blindly by 3 indepen- dent readers unaware of the treatment al- location and clinical events that occurred during the trial. Perfusion lung scans were reviewed and scored blindly by 2 indepen- dent readers according to the same proce- dure" Comment: review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes. Furthermore, the blinding of outcome as- sessment was ensured
Incomplete outcome data (attrition bias) All outcomes	Low risk	All data accounted for.
Selective reporting (reporting bias)	Low risk	Study reports data on all pre-specified out- comes.

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 57 thromboembolism (Review)

Meyer 1995 (Continued)

Other bias	Low risk	Study appears to be free from other sources of bias.	
Moreno-Palomares 2001			
Methods	Method of randomisation: not Concealment of allocation: no	Study design: randomised controlled trial. Method of randomisation: not stated. Concealment of allocation: not stated. Exclusions post-randomisation: none. Lost to follow-up: none.	
Participants	Age: mean 70 years LMWH, r Sex: LMWH 5 M/12 F, UFH Inclusion criteria: people with Exclusion criteria: people with		
Interventions	participant needed more than given over 12 hours Control: UFH: heparin sodiur Treatment duration: not stated	Treatment: LMWH: sodium dalteparin subcutaneously 200 U/kg over 24 hours. If the participant needed more than 180,000 U/day, the doses were divided into 2 and each given over 12 hours Control: UFH: heparin sodium 400 U/kg as an intravenous continuous infusion Treatment duration: not stated. Oral anticoagulation: oral dicocoumarol on 2nd day for 3 months	
Outcomes		Primary: progress of the Doppler. Secondary: post-phlebitic syndrome.	
Notes	Follow-up: 3, 6 and 12 month	Follow-up: 3, 6 and 12 months.	
Risk of bias			

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Insufficient information.
Allocation concealment (selection bias)	Unclear risk	Insufficient information.
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Insufficient information.
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Insufficient information.

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 58 thromboembolism (Review)

Moreno-Palomares 2001 (Continued)

Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Insufficient information.
Selective reporting (reporting bias)	Unclear risk	Insufficient information.
Other bias	Unclear risk	Insufficient information.
Ninet 1991		
Methods	Study design: randomised controlled trial. Method of randomisation: stratified to medical or surgical context in which VTE oc- curred Concealment of allocation: not blinded for treatment allocation, blinded for outcome assessment Exclusions post-randomisation: none. Lost to follow-up: 18 participants for assessment of change in thrombus size on venogram. No participants lost to follow-up for assessment of bleeding events	
Participants	Country: France (17 centres). Setting: hospital. No.: 166 participants undergoing medical or surgical procedures Age: estimated overall mean age 63 years. Sex: not stated. Inclusion criteria: recent (< 5 days) proximal DVT. Exclusion criteria: thrombosis affecting inferior vena cava; contraindication to heparin; platelets < 100,000/mm ³ ; blood disease; surgery < 3 days previously; contraindication for isotopic/venographic investigation; pulmonary vascular obstruction 30% or more (lung scan); 24 hours or more heparin or oral anticoagulant therapy; recent history (< 2 years) of cerebrovascular accident or thromboembolic episode; pregnancy	
Interventions	Treatment: LMWH: fraxiparine body weight-adjusted fixed dose (± 90 anti-factor Xa IU/kg, s.c., twice daily) Control: UFH: dose-adjusted APTT × 1.5 to 2.0, continuous i.v. infusion started with 20 IU/kg/hour. No bolus injection Treatment duration: 10 days. Oral anticoagulation: not defined for either group.	
Outcomes	Primary: change in thrombus size (Marder's score); recurrent venous thromboembolism (VTE) during initial treatment Secondary: haemorrhagic episodes during initial treatment; mortality at the end of fol- low-up	
Notes	Repeated venography on day 0 and day 10. Follow-up was not conducted prospectively at the study centre. 18 (8 versus 10) partic- ipants lost to follow-up. Follow-up by assessment on information noted and communicated by general practi- tioners	

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 59 thromboembolism (Review)

Ninet 1991 (Continued)

Risk of bias

Kisk of blas		
Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Insufficient information.
Allocation concealment (selection bias)	Unclear risk	Insufficient information.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Venography was evaluated blind by 2 inde- pendent radiologists (coded films)
Incomplete outcome data (attrition bias) All outcomes	High risk	Recurrences were excluded.
Selective reporting (reporting bias)	Low risk	Published report includes all expected out- comes.
Other bias	Low risk	There are more baseline risk factors in the UFH group compared to the CY 216 group. However, this difference was not sta- tistically significant

Prandoni 1992

Methods	Study design: randomised controlled trial. Method of randomisation: not stated. Concealment of allocation: not blinded for treatment allocation, blinded for outcome assessment Exclusions post-randomisation: none. Lost to follow-up: none.
Participants	Country: Italy. Setting: hospital. No.: 170 outpatients. Age: 86 (number over 65 years) Sex: 86 males. Inclusion criteria: proximal DVT. Exclusion criteria: clinically suspected PE at referral; episode of VTE in same leg within previous 2 years; anticoagulant treatment at referral; contraindication to heparin; preg- nancy; allergy to contrast material; residence far from hospital

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 60 thromboembolism (Review)

Prandoni 1992 (Continued)

Interventions	Treatment: LMWH: fraxiparine body weight-adjusted fixed dose; ± 90 anti-factor Xa IU/kg s.c., twice daily Control: UFH: dose-adjusted APTT × 1.5 to 2.0, continuous i.v. infusion started with 35,000 Units/24 hours. Initial bolus: 100 Units/kg i.v Treatment duration: at least 10 days; treatment cessation in INR > 2.0 Oral anticoagulation: Coumarin therapy initial dosage 5 mg started on day 7 of heparin treatment; INR 2.0 to 3.0
Outcomes	Primary: change in thrombus size (venogram day 1 and day 10); symptomatic recurrent DVT (including extension) or symptomatic PE; major haemorrhage during initial treat- ment Secondary: mortality; change in number of segmental defects on day 10 and day 0 lung scans
Notes	Follow-up: 6 months.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Participants were allocated treatment by a prescribed randomisation schedule
Allocation concealment (selection bias)	Low risk	Treatment was allocated by sealed envelopes.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	The venograms and perfusion lung scans of each participant were scored by a panel of 3 experienced observers who were unaware of treatment allocation and the sequence in which the tests were done (before or after treatment). All clinical end points were also reviewed by an adjudication committee un- aware of treatment allocation or other de- tails of participants
Incomplete outcome data (attrition bias) All outcomes	Low risk	No missing outcome data. No participant was lost to follow-up
Selective reporting (reporting bias)	Unclear risk	Insufficient information.

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 61 thromboembolism (Review)

Prandoni 1992 (Continued)

Other bias	Low risk	The study appears to be free of other sources of bias.	
Prandoni 2004			
Methods	Method of randomisation: com presented with DVT only or v Concealment of allocation: no assessment	Exclusions post-randomisation: none.	
Participants	than 3 weeks old) DVT of the of the following tests was requ the proximal vein system, echo clinical suspicion of DVT, vent scanning, and pulmonary ang presence of abnormal results of of PE was also accepted if a pe of PE when compared with th Exclusion criteria: age less that treatment, full-dose anticoagu more than 24 hours, haemody	Setting: hospital. No.: 720. Age: mean 66 years. Sex: 325 male, 395 female. Inclusion criteria: inpatients and outpatients with the clinical suspicion of an acute (less than 3 weeks old) DVT of the lower extremities and/or PE. A positive result of at least 1 of the following tests was required: ascending phlebography, compression ultrasound of the proximal vein system, echo colour Doppler scan of the calf vein system in the case of clinical suspicion of DVT, ventilation-perfusion scanning, spiral computed tomographic scanning, and pulmonary angiography in the case of clinical suspicion of PE. In the presence of abnormal results of an ultrasound test of the lower extremities, the diagnosis of PE was also accepted if a perfusion lung scan was compatible with a high probability of PE when compared with the chest x-ray Exclusion criteria: age less than 18 years, pregnancy, contraindications to anticoagulant treatment, full-dose anticoagulant treatment (either heparin or oral anticoagulants) for more than 24 hours, haemodynamic instability, previous (less than 1 year earlier) episode of VTE, life expectancy less than 3 months, poor compliance, and geographic inacces-	
Interventions	85 IU/kg twice daily Control: UFH: an i.v. bolus of doses adjusted to body weight less than 50 kg; 5000 IU plus kg; and 6000 IU plus 17,500 II The first APTT was measured first 48 hours were scheduled was managed on the basis of d Treatment duration: At least 5 days	Control: UFH: an i.v. bolus of heparin sodium and a s.c. injection of heparin calcium in doses adjusted to body weight (4000 IU i.v. plus 12500 IU s.c. in participants weighing less than 50 kg; 5000 IU plus 15,000 IU, respectively, in participants weighing 50 to 70 kg; and 6000 IU plus 17,500 IU, respectively, in participants weighing more than 70 kg). The first APTT was measured after 6 hours, and subsequent dose adjustments during the first 48 hours were scheduled twice daily. After the first 48 hours, UHF administration was managed on the basis of daily APTT determinations Treatment duration: At least 5 days; heparin cessation if INR was > 2.0 for 2 consecutive days Oral anticoagulation: warfarin sodium was started within the first 2 days and continued	

Prandoni 2004 (Continued)

Outcomes	Primary: recurrent thromboembolism and mortality during heparin treatment and fol- low-up Secondary: Major bleeding during the period of heparin treatment and the subsequent 48 hours
Notes	Follow-up: 3 months.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Randomisation was performed with a com- puter algorithm.
Allocation concealment (selection bias)	Low risk	Central allocation by a 24-hour telephone service.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Information on all suspected outcome events and deaths was reviewed and clas- sified by a central adjudication committee blinded to treatment assignment
Incomplete outcome data (attrition bias) All outcomes	Low risk	No missing outcome data.
Selective reporting (reporting bias)	Low risk	Published report includes all expected out- comes.
Other bias	Low risk	The study appears to be free of other sources of bias.

Pérez de Llano 2003

Methods	Study design: multicentre, prospective open study Method of randomisation: SAS statistics computer program Concealment of allocation: none Exclusions post-randomisation: none Lost to follow-up: none
Participants	Country: Spain. Setting: 3 hospitals. No.: enoxaparin 29, UFH 21

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 63 thromboembolism (Review)

Pérez de Llano 2003 (Continued)

	Age: enoxaparin mean 66.5 \pm 16.2 years, UFH mean 65.9 \pm 16.3 years Sex: enoxaparin 20 M/9 F, UFH 14 M/ 7 F Inclusion criteria: people diagnosed with pulmonary thromboembolism (PTE) diag- nosed by ventilation-perfusion scan or plethysmography Exclusion criteria: people with a previous DVT, PTE with haemodynamic repercussion, known factor of hypercoagulability, anticoagulant treatment, pregnancy, formal consid- eration for anticoagulation or serious concomitant illnesses
Interventions	Treatment: enoxaparin 1 mg/kg weight every 12 hours. Control: 5% sodium heparin 5000 IU initial bolus through an infusion pump adjusted to the partial thromboplastin time results to an approximated dose of 35,000 IU/day Treatment duration: until a target INR of 2 to 3 was reached Oral anticoagulation: acenocoumarol.
Outcomes	Primary: recurrence of DVT (if plethysmography showed a new venous region affected, if there was a proximal thrombus extension > 5 cm or if arteriography showed new intra- luminal defects) or PE (if perfusion scan showed perfusion defects that had not existed in the initial exploration) and major bleeding (intracranial, retroperitoneal, requiring transfusion or haemoglobin < 2 or more points)
Notes	Follow-up: 1, 3 and 6 months.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "Patients were randomised from the lists of enrolled patients at each centre using the SAS statistics program" Comment: low risk of bias.
Allocation concealment (selection bias)	Unclear risk	Method of allocation concealment not stated. Comment: insufficient information to per- mit judgement of low or high risk
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	It is not stated whether the outcome asses- sors were blinded to treatment and there- fore the risk of bias was deemed unclear
Incomplete outcome data (attrition bias) All outcomes	Low risk	All data accounted for.

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 64 thromboembolism (Review)

Pérez de Llano 2003 (Continued)

Selective reporting (reporting bias)	High risk	Study authors discuss the length of hospital stay but it was not a prespecified outcome
Other bias	Low risk	Study appears to be free from other sources of bias.
Riess 2003		
Methods	Study design: randomised controlled trial. Method of randomisation: not stated. Concealment of allocation: not blinded for treatment allocation, blinded for outcome assessment Exclusions post-randomisation: 92 participants. Lost to follow-up: 22 participants.	
Participants	Country: 121 centres in Germany and the Czech Republic. Setting: hospital and out of hospital. No.: 1220 participants. Age: mean 61 years. Sex: 677 males. Inclusion criteria: men older than 18 years with objectively confirmed acute proximal DVT for fewer than 3 weeks after given written informed consent Exclusion criteria: isolated calf vein thrombosis; planned fibrinolysis or operation; clini- cally severe PE; heparin application within 8 days of enrolment (except treatment in the past 24 hours), treatment with VKA for > 24 hours before start of study medication; hypertension with systolic blood pressure > 200 mmHg and diastolic blood pressure > 105 mmHg; known malignant tumour as known cause for the venous occlusion; severe renal or hepatic insufficiency; surgery of the head, chest or abdomen in the past 8 days; intervention in the central nervous system in the past 14 days; evident disseminated intravascular coagulation; clinical condition with an increased risk of bleeding compli- cations during the treatment time; gastrointestinal bleeding or gastric ulcer in the past 4 weeks; contraindication against VKA or known intolerability against heparin; platelet count < 100,000/µL; pregnancy, treatment with platelet inhibitors	
Interventions	Treatment: LMWH: certoparin fixed unadjusted dose 8000 anti-factor Xa IU s.c., b.d. for 10 to 14 days Control: UFH: initial bolus i.v. of 5000 IU followed by continuous infusion starting dose of 20 IU/kg/hour of an adjusted dose UFH to maintain an APTT of 1.5 to 2.5 × the control value	
Outcomes	Primary: incidence of VTE at the end of fo Secondary: incidence of recurrent VTE an mortality at the end of follow-up	llow-up. Id major bleeding during initial treatment;
Notes	Follow-up: 6 months.	
Risk of bias		

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 65 thromboembolism (Review)

Riess 2003 (Continued)

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Insufficient information about sequence generation.
Allocation concealment (selection bias)	Low risk	Randomisation was carried out using a cen- tral telephone system. The assignment to 1 of the treatment groups was documented and could not be changed afterwards
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	All events were evaluated by an indepen- dent end point committee blinded for treatment groups
Incomplete outcome data (attrition bias) All outcomes	Low risk	An intent-to-treat analysis confirmed the results of the primary 'per protocol' analysis
Selective reporting (reporting bias)	Low risk	Published report includes all expected out- comes.
Other bias	Low risk	The study appears to be free of other sources of bias.

Simonneau 1993

Methods	 Study design: randomised controlled trial. Method of randomisation: treatment assignments: sealed envelopes, block randomisation using standard random number table and sealed envelopes Concealment of allocation: not blinded for treatment allocation; blinded for outcome assessment Exclusion post-randomisation: 1 (distal DVT). Lost to follow-up: for qualitative and quantitative venogram assessment: 17 participants lost to follow-up (treatment cessation before day 10 (5 participants); exclusion post randomisation (1 participant); unassessable venograms due to technical problems (11 participants))
Participants	Country: 16 European centres. Setting: hospital. No.: 134 participants. Age: Mean 63 years. Sex: 73 males. Inclusion criteria: proximal DVT with or without suspected PE, but with symptoms <

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 66 thromboembolism (Review)

Simonneau 1993 (Continued)

	5 days Exclusion criteria: active bleeding or disorders contraindicating anticoagulant therapy; surgery in previous 7 days; pregnancy; aspirin, ticlopidine, sulfinpyrazone or non- steroidal anti-inflammatory treatment within 7 days before study entry; associated severe PE requiring thrombolytic therapy or surgery; use of curative heparin therapy for > 24 hours or > 25,000 Units of heparin during 24 hours before referral; previous implanta- tion of vena cava filter
Interventions	Treatment: LMWH: enoxaparin, clexane body weight-adjusted fixed dose (1 mg/kg ± 100 anti-factor Xa IU/kg, s.c., twice daily) Control: UFH: dose-adjusted APTT × 1.5 to 2.5, continuous i.v. infusion started with 500 Units/kg/24 hours (25,000 Units/5 mL in saline) Treatment duration: 10 days. Oral anticoagulation: started on day 10 for at least 3 months; INR 2.0 to 3.0
Outcomes	Primary: change in thrombus size (quantitative venographic score, Marder) between day 0 and day 10; recurrent VTE during 10 days of treatment (asymptomatic and symptomatic DVT and PE); major bleeding during 10 days of treatment Secondary: minor bleeding; follow-up at 3 months to record VTE recurrence, bleeding and deaths; qualitative assessment of venogram evolution between day 0 and day 10
Notes	Repeated venography on day 10. Follow-up: 3 months.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	The randomisation code was drafted by means of a standard random number table randomising in blocks of 4
Allocation concealment (selection bias)	Low risk	The participants' treatment assignments were taken from sealed envelopes
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Venograms, perfusion lung scans and pul- monary angiograms were subsequently re- viewed by a central independent panel of 2 consultant specialists unaware of the treat- ment allocation
Incomplete outcome data (attrition bias) All outcomes	Low risk	Reasons for missing outcome data unlikely to be related to true outcome

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 67 thromboembolism (Review)

Simonneau 1993 (Continued)

Selective reporting (reporting bias)	Low risk	Published report includes all expected out- comes.	
Other bias	Low risk	The study appears to be free of other sources of bias.	
Simonneau 1997			
Methods	Study design: randomised controlled trial. Method of randomisation: centrally controlled, computerised. Concealment of allocation: not blinded for treatment allocation, blinded for outcome assessment Exclusions post-randomisation: none. Lost to follow-up: none.		
Participants	 Country: France, Belgium and Switzerland. Setting: hospital. No.: 612 participants. Age: mean 67 years. Sex: 172 males. Inclusion criteria: clinically suspected acute PE. PE objectively documented by pulmonary angiography or ventilation-perfusion lung scanning indicating a high probability of PE or showing indeterminate results but accompanied by DVT confirmed by venography or compression ultrasonography Exclusion criteria: massive PE requiring thrombolytic therapy or pulmonary embolectomy; active bleeding or disorders contraindicating anticoagulant therapy; anticoagulant therapy at a therapeutic dose for > 24 hours; life expectancy < 3 months; severe hepatic or renal failure; likely non-compliance; pregnancy 		
Interventions	Treatment: LMWH: tinzaparin, innohep in body weight-adjusted fixed dose, s.c., o.d Control: UFH: APTT-adjusted dose, continuous i.v. infusion after an initial i.v. bolus of 50 IU/kg Treatment duration: at least 5 days; treatment cessation if INR was 2.0 or above on 2 measurements made 24 hours apart Oral anticoagulation: started between the first and third days of initial treatment and continued for at least 3 months; INR 2.0 to 3.0		
Outcomes	Primary: symptomatic recurrent VTE during initial treatment (8 days) and at the end of follow-up (day 90); major haemorrhage during initial treatment (8 days) and at the end of follow-up (day 90); death at end of follow-up (day 90)		
Notes	Follow-up: 90 days. 1 participant allocated to UFH and 3 participants allocated to LMWH did not receive the study drug		
Risk of bias			
Bias	Authors' judgement	Support for judgement	

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 68 thromboembolism (Review)

Simonneau 1997 (Continued)

Random sequence generation (selection bias)	Low risk	Central randomisation was performed with the use of a 24-hour computer service
Allocation concealment (selection bias)	Low risk	Central randomisation was performed with the use of a 24-hour computer service
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Data on all potential outcome events were submitted to an independent adjudication committee whose members were unaware of the treatment assignments
Incomplete outcome data (attrition bias) All outcomes	Low risk	An intention-to-treat analysis was per- formed, but the authors do not give any in- formation about loss to follow-up
Selective reporting (reporting bias)	Low risk	Published report includes all expected out- comes.
Other bias	Low risk	The study appears to be free of other sources of bias.

Thery 1992

Methods	Study design: dose-finding controlled, randomised trial. Method of randomisation: not stated Concealment of allocation: not blinded for treatment allocation, blinded for outcome assessment Exclusions post-randomisation: none. Lost to follow-up: none.
Participants	Country: France, Setting: hospital. No.: 68: Fraxparine 35, UFH 33 Age: Fraxiparine mean 60.1 (SD 2.9) years, UFH mean 64.2 (SD 2.5) years Sex: Fraxiparine 17 M/18 F, UFH 14 M/19 F Inclusion criteria: adults > 18 years with a recent angiographically proved PE (within 3 days of the onset of symptoms) and with a pulmonary vascular obstruction assessed by the local radiologists between 15% and 55% (index of severity according to Miller 5 to 18) Exclusion criteria: angiographically determined vascular obstruction < 15% or > 55%, any sign of clinical severity defined as shock, acute cor pulmonale or right heart fail- ure, any contraindication to heparin, active peptic ulcer, recent history of cerebrovascu-

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 69 thromboembolism (Review)

Thery 1992 (Continued)

	lar haemorrhage or ischaemia, known bleeding tendency, previous history of heparin- induced thrombocytopenia, haemorrhagic diathesis, pre-existing coagulation disorders, severe renal or hepatic dysfunction, severe systemic hypertension, known pericarditis or endocarditis, pregnancy, pre-existing DVT or PE within 12 months preceding the inclusion or use of thrombolytic agents, heparin at therapeutic doses for more than 48 hours before inclusion, oral anticoagulants, acetylsalicylic acid or ticlopidine during the 7 days before inclusion, any contraindication to isotopic or angiographic investigations and free-floating inferior vena cava thrombus
Interventions	Treatment: LMWH: Fraxiparine 400 anti-factor Xa IU U/kg in 2 daily injections Control: UFH: i.v. bolus injection of 50 IU/kg followed by continuous infusion of an initial dose of 600 IU/kg Treatment duration: 14 days Oral anticoagulation: none
Outcomes	Primary: pulmonary vascular obstruction Secondary: clinical recurrence of VTE, death and haemorrhagic complications
Notes	Follow-up: 8 days Before completion of the trial, enrolment in 2 Fraxiparine groups stopped because of a high incidence of major bleedings. Those 2 groups were given Fraxiparine at a high dose of 600 and 900 anti-factor Xa IU/kg. Data from these groups were not included in the analyses in this review

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Quote: "Random treatment allocation schedules were prepared for each clinical centre using sealed treatment allocation en- velopes" Comment: insufficient information about the sequence generation process to permit judgement of 'Low risk' or 'High risk'
Allocation concealment (selection bias)	Unclear risk	Quote: "Random treatment allocation schedules were prepared for each clinical centre using sealed treatment allocation en- velopes" Comment: although the use of assignment envelopes is described, it remains unclear whether envelopes were sequentially num- bered and opaque
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Not blinded for treatment allocation. Comment: given the clinical outcomes of the study, review authors judged that the non-blinding of the participants and staff

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 70 thromboembolism (Review)

Thery 1992 (Continued)

		was unlikely to have affected the outcomes
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "the study could not be per- formed double-blind because of the dif- ferent modes of administration and above all the need for dosage adjustments in the UFH group. However, the main assessment criterion was blindly evaluated by a central independent panel of three radiologists" Comment: review authors judged that the non-blinding of the participants and staff was unlikely to have affected the outcomes. Furthermore, the blinding of outcome as- sessment was ensured
Incomplete outcome data (attrition bias) All outcomes	High risk	Reasons for losses to follow-up not clearly stated.
Selective reporting (reporting bias)	Low risk	Study reports data on all pre-specified out- comes.
Other bias	Low risk	Study appears to be free from other sources of bias.

APTT: activated partial thromboplastin time

cm: centimetre DVT: deep vein thrombosis F: female INR: International normalised ratio IU: International units i.v.: intravenous kg: kilogram LMWH: low molecular weight heparin M: male mg: milligram mL: millilitre mm: millimetre mmHg: millimetres of mercury NSAID: nonsteroidal anti-inflammatory drug PE: pulmonary embolism PTE: pulmonary thromboembolism o.d.: once daily s.c.: subcutaneous SD: standard deviation UFH: unfractionated heparin VKA: vitamin K antagonists VTE: venous thromboembolism

Characteristics of excluded studies [ordered by study ID]

Study	Reason for exclusion
Aiach 1989	Low molecular weight heparin dosage was adjusted.
Albada 1989	The results from participants treated for venous thrombosis of the upper limb and for pulmonary embolism could not be distinguished from those of people with leg vein thrombosis, and the outcome was incompletely evaluated
Banga 1993	This was a dose-finding study.
Bratt 1985	Intravenous route of administration of low molecular weight heparin, and adjustments were made to dose for this treatment
Bratt 1990	Low molecular weight heparin dosage was adjusted.
de Valk 1995	This was a dose-finding study.
Handeland 1990	This was a dose-finding study.
Harenberg 1989	Abstract with incomplete data.
Harenberg 1990	The results from people treated for venous thrombosis of the upper limb and for pulmonary embolism could not be distinguished from those of participants with leg vein thrombosis, and the outcome was incompletely evaluated
Harenberg 2000b	Abstract with incomplete data.
Holm 1986	Low molecular weight heparin dosage was adjusted.
Kearon 2006	The administration of unfractionated heparin was not in adjusted dose
Lockner 1985	Intravenous route of administration of low molecular weight heparin
Lockner 1986	Intravenous route of administration of low molecular weight heparin
Ly 1985	Adjustment of low molecular weight heparin dosages.
Monreal 1993	The 2 treatment strategies differ with respect to long-term treatment
Monreal 1994	The 2 treatment strategies differ with respect to long-term treatment
Notarbartolo 1988	Dosage of unfractionated heparin was not adjusted.
Quiros 2001	Not a randomised controlled trial.
Riess 2014	Substudy of Harenberg 1990 and Reiss 2003 studies which are already included in the review

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 72 thromboembolism (Review)

(Continued)

Siguret 2011	Comparison group were not treated with unfractionated heparin
Stricker 1999	The main outcome of the study was the effect on haemostatic markers which is not within the scope of our review
Tedoldi 1993	Dosage of unfractionated heparin was not adjusted.
Ucar 2015	Participants were given thrombolytic treatment.
Vogel 1987	Intravenous route of administration of low molecular weight heparin
Zanghi 1988	Dosage of unfractionated heparin was not adjusted.

Characteristics of ongoing studies [ordered by study ID]

NCT00796692

Trial name or title	Nadroparin for the Initial Treatment of Pulmonary Thromboembolism (NATSPUTE)
Methods	Multicentre, randomised, open-label, parallel assignment controlled trial
Participants	Inclusion criteria: 18 to 75 years of age, symptomatic non-massive PTE confirmed either by high probability ventilation-perfusion lung scanning (V/Q scan) or by the presence of intraluminal filling defect on spiral computed tomographic pulmonary angiography (CTPA), haemodynamically stabile, anatomic obstruction no more than 2 lobes on CTPA, or defect no more than 7 segments on V/Q scan, and normal right ventricular function, symptoms within 15 days, written informed consent obtained before randomisation Exclusion criteria: unfractionated heparin anticoagulation for more than 36 hours prior enrolment, massive PTE or sub-massive PTE requiring thrombolytic therapy or pulmonary embolectomy, active bleeding or disorders contraindicating anticoagulant therapy, chronic thromboembolism pulmonary hypertension (CTEPH) without evidence of recent episode, severe hepatic or renal failure, allergy to heparin, other components of tinzaparin or acenocoumarol, pregnant status, a life expectancy of less than 3 months, previous thrombocytopaenia induced by heparin, thrombocytopaenia < 100,000/mm ³
Interventions	Treatment: LMWH given with a weight-adjusted dose of 86 international anti-factor Xa units of nadroparin (Fraxiparine) per kilogram of body weight (86 anti-factor Xa IU/kg) subcutaneously every 12 hours, which will be used at least 5 to 7 days Control: UFH is received with an initial bolus dose of 80 IU per kilogram, followed by a continuous intravenous infusion at an initial rate of 18 IU per kilogram per hour. The dose is subsequently adjusted so that the activated partial thromboplastin time (APTT) would be 1.5 to 2.5 times the control value in normal subjects. The tests are performed 4 hours after the start of treatment, whenever a sub-therapeutic APTT had been measured after a dose adjustment, and otherwise daily. UFH will be used at least 5 to 7 days Treatment duration: 5 to 7 days. Oral anticoagulation: warfarin.
Outcomes	Primary: clinical and image (including V/Q scan and CTPA) improvement at 14 days Secondary: recurrent venous thromboembolism (VTE), major bleeding, death and heparin-induced throm- bocytopaenia at 3 months

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 73 thromboembolism (Review)

NCT00796692 (Continued)

Starting date	June 2002
Contact information	Professor Chen Wang, Beijing Institute of Respiratory Medicine, Beijing Chao Yang Hospital, China
Notes	Study authors have been contacted for further information but no response received to date

IU: international units kg: kilogram LMWH: low molecular weight heparin mm: millimetre PTE: pulmonary thromboembolism UFH: unfractionated heparin

DATA AND ANALYSES

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Incidence of recurrent venous thromboembolism during initial treatment	18	6238	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.69 [0.49, 0.98]
1.1 Nadroparin versus unfractionated heparin	5	950	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.50 [0.20, 1.26]
1.2 Tinzaparin versus unfractionated heparin	1	612	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.52 [0.26, 8.80]
1.3 Enoxaparin versus unfractionated heparin	5	1143	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.51 [0.27, 0.98]
1.4 Dalteparin versus unfractionated heparin	3	495	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.08 [0.07, 17.43]
1.5 Reviparin versus unfractionated heparin	1	1021	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.07 [0.52, 2.19]
1.6 Certoparin versus unfractionated heparin	3	2017	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.68 [0.35, 1.32]
2 Incidence of recurrent venous thromboembolism at the end of follow-up	22	9489	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.72 [0.59, 0.88]
2.1 Nadroparin versus unfractionated heparin	5	1730	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.74 [0.49, 1.11]
2.2 Tinzaparin versus unfractionated heparin	3	1581	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.76 [0.41, 1.40]
2.3 Ardeparin versus unfractionated heparin	1	80	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.13 [0.00, 6.49]
2.4 Enoxaparin versus unfractionated heparin	6	2043	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.76 [0.49, 1.17]
2.5 Dalteparin versus unfractionated heparin	2	264	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.71 [0.42, 7.02]
2.6 Reviparin versus unfractionated heparin	2	1784	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.68 [0.44, 1.05]
2.7 Certoparin versus unfractionated heparin	3	2007	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.63 [0.40, 0.99]
3 Incidence of recurrent venous thromboembolism at 1 month follow-up	4	1741	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.90 [0.56, 1.44]
3.1 Nadroparin versus unfractionated heparin	1	170	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.50 [0.10, 2.55]
3.2 Enoxaparin versus unfractionated heparin	2	550	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.59 [0.24, 1.48]
3.3 Reviparin versus unfractionated heparin	1	1021	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.15 [0.64, 2.06]

Comparison 1. LMWH versus UFH in patients with venous thromboembolism

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 75 thromboembolism (Review)

4 Incidence of recurrent venous thromboembolism at 3 months' follow-up	16	6661	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.71 [0.56, 0.90]
4.1 Nadroparin versus unfractionated heparin	5	1730	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.77 [0.49, 1.21]
4.2 Tinzaparin versus unfractionated heparin	2	1044	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.54 [0.26, 1.08]
4.3 Enoxaparin versus unfractionated heparin	6	2043	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.76 [0.49, 1.17]
4.4 Dalteparin versus unfractionated heparin	1	60	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.0 [0.0, 0.0]
4.5 Reviparin versus unfractionated heparin	2	1784	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.68 [0.44, 1.05]
5 Incidence of recurrent venous thromboembolism at 6 months' follow-up	7	2841	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.68 [0.48, 0.96]
5.1 Nadroparin versus unfractionated heparin	2	570	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.66 [0.37, 1.19]
5.2 Dalteparin versus unfractionated heparin	1	204	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.71 [0.42, 7.02]
5.3 Certoparin versus	3	2007	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.63 [0.40, 0.99]
unfractionated heparin				
5.4 Enoxaparin versus unfractionated heparin	1	60	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.0 [0.0, 0.0]
6 Reduction in thrombus size (pre- and post-treatment venograms)	16	2909	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.71 [0.61, 0.82]
6.1 Nadroparin versus unfractionated heparin	4	507	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.72 [0.50, 1.05]
6.2 Ardeparin versus unfractionated heparin	1	75	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.37 [0.14, 0.99]
6.3 Enoxaparin versus unfractionated heparin	1	117	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.34 [0.17, 0.71]
6.4 Dalteparin versus unfractionated heparin	5	650	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.25 [0.90, 1.73]
6.5 CY 222 versus unfractionated heparin	1	59	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.91 [0.32, 2.62]
6.6 Reviparin versus unfractionated heparin	1	649	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.59 [0.43, 0.80]
6.7 Certoparin versus unfractionated heparin	2	649	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.70 [0.50, 0.98]
6.8 Bemiparin versus unfractionated heparin	1	203	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.42 [0.24, 0.74]
7 Incidence of major haemorrhagic episodes (during initial treatment)	25	8780	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.69 [0.50, 0.95]
7.1 Nadroparin versus unfractionated heparin	7	1964	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.44 [0.19, 1.01]
7.2 Tinzaparin versus unfractionated heparin	3	1581	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.71 [0.41, 1.22]
7.3 Enoxaparin versus unfractionated heparin	5	1143	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.22 [0.54, 2.75]

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 76 thromboembolism (Review)

7.4 Dalteparin versus	4	765	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.15 [0.02, 1.44]
unfractionated heparin				
7.5 CY 222 versus	1	68	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.13 [0.01, 1.34]
unfractionated heparin				
7.6 Reviparin versus	1	1021	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.26 [0.49, 3.19]
unfractionated heparin				
7.7 Certoparin versus	3	2017	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.50 [0.25, 1.00]
unfractionated heparin				
7.8 Bemiparin versus	1	221	Peto Odds Ratio (Peto, Fixed, 95% CI)	7.39 [0.46, 118.89]
unfractionated heparin				
8 Overall mortality at the end of	24	9663	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.84 [0.70, 1.01]
follow-up				
8.1 Nadroparin versus	5	1504	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.77 [0.48, 1.22]
unfractionated heparin				
8.2 Tinzaparin versus	3	1581	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.03 [0.69, 1.53]
unfractionated heparin				
8.3 Ardeparin versus	1	80	Peto Odds Ratio (Peto, Fixed, 95% CI)	$0.0 \ [0.0, \ 0.0]$
unfractionated heparin				
8.4 Enoxaparin versus	6	2043	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.92 [0.64, 1.31]
unfractionated heparin				
8.5 Dalteparin versus	3	490	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.46 [0.13, 1.60]
unfractionated heparin				
8.6 Reviparin versus	2	1784	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.89 [0.59, 1.35]
unfractionated heparin				
8.7 Certoparin versus	3	2007	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.59 [0.36, 0.97]
unfractionated heparin				
8.8 Bemiparin versus	1	174	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.95 [0.13, 6.90]
unfractionated heparin				

Comparison 2. LMWH versus UFH in patients with proximal deep venous thrombosis

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Incidence of recurrent venous thromboembolism at the end of follow-up	10	4672	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.57 [0.44, 0.75]
1.1 Nadroparin versus unfractionated heparin	3	864	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.72 [0.44, 1.19]
1.2 Tinzaparin versus unfractionated heparin	1	432	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.42 [0.17, 1.01]
1.3 Enoxaparin versus unfractionated heparin	2	634	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.65 [0.32, 1.32]
1.4 Reviparin versus unfractionated heparin	1	763	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.31 [0.15, 0.63]
1.5 Certoparin versus unfractionated heparin	2	1758	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.64 [0.40, 1.03]

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 77 thromboembolism (Review)

venous thrombosis at the end of follow-up	0, 0.0] 0.41, 0.91] 0.41, 1.43] 0.12, 1.16] 0.34, 1.63]
2 Incidence of recurrent deep 7 2681 Peto Odds Ratio (Peto, Fixed, 95% CI) 0.61 [0] venous thrombosis at the end of follow-up 2 765 Peto Odds Ratio (Peto, Fixed, 95% CI) 0.76 [0]	0.41, 1.43] 0.12, 1.16]
venous thrombosis at the end of follow-up 2.1 Nadroparin versus 3 765 Peto Odds Ratio (Peto, Fixed, 95% CI) 0.76 [0	0.41, 1.43] 0.12, 1.16]
of follow-up2.1 Nadroparin versus3765Peto Odds Ratio (Peto, Fixed, 95% CI)0.76 [0]).12, 1.16]
1).12, 1.16]
unfractionated heparin	
2.2 Tinzaparin versus1432Peto Odds Ratio (Peto, Fixed, 95% CI)0.37 [0	.34, 1.63]
unfractionated heparin	0.34, 1.63]
•	
unfractionated heparin	
•).17, 1.21]
unfractionated heparin 2.5 Bemiparin versus 1 221 Peto Odds Ratio (Peto, Fixed, 95% CI) 0.13 [0	0.01, 2.14]
unfractionated heparin	.01, 2.14]
-	0.28, 0.74]
embolism at the end of	
follow-up	
*	0.27, 1.60]
unfractionated heparin	
*).14, 1.95]
unfractionated heparin	05 5 071
3.3 Enoxaparin versus1500Peto Odds Ratio (Peto, Fixed, 95% CI)0.52 [0unfractionated heparin	0.05, 5.07]
).10, 0.73]
unfractionated heparin	.10, 0.75]
-).11, 0.92]
unfractionated heparin	
•	0.20, 18.86]
unfractionated heparin	
•	0.27, 0.80]
and post-treatment venograms)	2607/1
4.1 Bemiparin versus1203Peto Odds Ratio (Peto, Fixed, 95% CI)0.42 [0unfractionated heparin	0.24, 0.74]
).22, 9.90]
unfractionated heparin	.22, 7.90]
-).29, 0.85]
episodes (during initial	-
treatment)	
	0.09, 1.85]
unfractionated heparin	
5.2 Tinzaparin versus1432Peto Odds Ratio (Peto, Fixed, 95% CI)0.19 [0unfractionated heparin	0.06, 0.59]
•	0.42, 6.87]
unfractionated heparin	.42, 0.07]
	0.26, 1.17]
unfractionated heparin	
	0.47, 0.85]
follow-up	
	0.38, 1.24]
unfractionated heparin	

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 78 thromboembolism (Review)

6.2 Tinzaparin versus unfractionated heparin	1	432	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.48 [0.23, 1.00]
6.3 Enoxaparin versus	2	634	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.74 [0.37, 1.50]
unfractionated heparin				
6.4 Reviparin versus	1	763	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.79 [0.32, 1.91]
unfractionated heparin				
6.5 Certoparin versus	2	1758	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.54 [0.30, 0.96]
unfractionated heparin				
6.6 Bemiparin versus	1	174	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.95 [0.13, 6.90]
unfractionated heparin				

Comparison 3. LMWH versus UFH in patients with pulmonary embolism

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Incidence of recurrent venous thromboembolism at the end of follow-up	7	1407	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.90 [0.50, 1.61]
1.1 Tinzaparin versus unfractionated heparin	2	680	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.84 [0.26, 2.77]
1.2 Enoxaparin versus unfractionated heparin	3	396	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.89 [0.37, 2.16]
1.3 Dalteparin versus unfractionated heparin	1	60	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.0 [0.0, 0.0]
1.4 Reviparin versus unfractionated heparin	1	271	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.96 [0.35, 2.64]
2 Reduction in thrombus size (pre- and post-treatment venograms)	2	106	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.36 [0.23, 8.16]
2.1 Nadroparin versus unfractionated heparin	1	52	Peto Odds Ratio (Peto, Fixed, 95% CI)	5.53 [0.32, 95.93]
2.2 Dalteparin versus unfractionated heparin	1	54	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.54 [0.05, 5.43]
3 Mean change in pulmonary vascular obstruction severity score	2	106	Mean Difference (IV, Fixed, 95% CI)	-3.14 [-4.39, -1.90]
3.1 Nadroparin versus unfractionated heparin	1	52	Mean Difference (IV, Fixed, 95% CI)	-3.28 [-4.55, -2.01]
3.2 Dalteparin versus unfractionated heparin	1	54	Mean Difference (IV, Fixed, 95% CI)	1.0 [-5.94, 7.94]
4 Incidence of major haemorrhagic episodes (during initial treatment)	3	178	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.44 [0.04, 4.29]
4.1 Nadroparin versus unfractionated heparin	1	68	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.12 [0.01, 2.02]
4.2 Enoxaparin versus unfractionated heparin	1	50	Peto Odds Ratio (Peto, Fixed, 95% CI)	5.61 [0.11, 297.44]

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 79 thromboembolism (Review)

4.3 Dalteparin versus unfractionated heparin	1	60	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.0 [0.0, 0.0]
5 Overall mortality at end of follow-up	3	178	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.70 [0.17, 16.71]
5.1 Nadroparin versus unfractionated heparin	1	68	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.94 [0.06, 15.40]
5.2 Enoxaparin versus unfractionated heparin	1	50	Peto Odds Ratio (Peto, Fixed, 95% CI)	5.61 [0.11, 297.44]
5.3 Dalteparin versus unfractionated heparin	1	60	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.0 [0.0, 0.0]

Comparison 4. LMWH versus UFH in patients with venous thromboembolism and malignant disease

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Mortality at the end of follow-up	6		Peto Odds Ratio (Peto, Fixed, 95% CI)	Subtotals only
1.1 Mortality in patients with malignant disease	6	446	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.53 [0.33, 0.85]
1.2 Mortality in patients with malignant disease in trial with adequate concealment	5	430	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.51 [0.31, 0.82]

Comparison 5. LMWH versus UFH in patients with venous thromboembolism without malignant disease

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Mortality at the end of follow-up	6		Peto Odds Ratio (Peto, Fixed, 95% CI)	Subtotals only
1.1 Mortality in patients without malignant disease	6	2139	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.97 [0.61, 1.56]
1.2 Mortality in patients without malignant disease in trials with adequate concealment	5	1951	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.00 [0.62, 1.62]

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 80 thromboembolism (Review)

Comparison 6. LMWH versus subcutaneous UFH in patients with venous thromboembolism	Comparison 6.	LMWH versus subcutaneous UFH	I in patients with ven	ous thromboembolism
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Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Incidence of recurrent venous thromboembolism at the end of follow-up	3	1403	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.05 [0.56, 1.95]
2 Incidence of major haemorrhagic episodes (during initial treatment)	4	1471	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.91 [0.50, 1.67]
3 Overall mortality at the end of follow-up	3	1403	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.46 [0.91, 2.35]

Comparison 7. LMWH versus intravenous UFH in patients with venous thromboembolism

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Incidence of recurrent venous thromboembolism at the end of follow-up	21	8375	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.69 [0.56, 0.86]
2 Incidence of major haemorrhagic episodes (during initial treatment)	21	7309	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.62 [0.43, 0.90]
3 Overall mortality at the end of follow-up	21	8260	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.77 [0.63, 0.93]

Comparison 8. LMWH versus UFH: all randomised controlled trials with adequate concealment of allocation

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Incidence of recurrent venous thromboembolism during initial treatment	10	4862	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.72 [0.50, 1.05]
1.1 Nadroparin versus unfractionated heparin	3	716	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.50 [0.18, 1.39]
1.2 Tinzaparin versus unfractionated heparin	1	612	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.52 [0.26, 8.80]
1.3 Enoxaparin versus unfractionated heparin	3	1034	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.48 [0.24, 0.96]
1.4 Reviparin versus unfractionated heparin	1	1021	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.07 [0.52, 2.19]
1.5 Certoparin versus unfractionated heparin	2	1479	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.79 [0.40, 1.58]

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 81 thromboembolism (Review)

2 Incidence of recurrent venous thromboembolism at the end of follow-up	14	6984	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.76 [0.60, 0.96]
2.1 Nadroparin versus unfractionated heparin	4	1436	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.70 [0.45, 1.10]
2.2 Tinzaparin versus unfractionated heparin	2	1044	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.54 [0.26, 1.08]
2.3 Ardeparin versus unfractionated heparin	1	81	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.13 [0.00, 6.65]
2.4 Enoxaparin versus unfractionated heparin	4	1934	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.79 [0.51, 1.22]
2.5 Reviparin versus unfractionated heparin	1	1020	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.08 [0.62, 1.89]
2.6 Certoparin versus unfractionated heparin	2	1469	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.73 [0.42, 1.25]
3 Incidence of recurrent venous thromboembolism at 3 months'	11	5435	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.79 [0.60, 1.02]
follow-up				
3.1 Nadroparin versus unfractionated heparin	4	1436	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.73 [0.44, 1.22]
3.2 Tinzaparin versus unfractionated heparin	2	1044	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.54 [0.26, 1.08]
3.3 Enoxaparin versus unfractionated heparin	4	1934	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.79 [0.51, 1.22]
3.4 Reviparin versus unfractionated heparin	1	1021	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.09 [0.62, 1.90]
4 Reduction in thrombus size (pre- and post-treatment venograms)	5	753	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.49 [0.37, 0.66]
4.1 Nadroparin versus unfractionated heparin	2	302	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.49 [0.31, 0.77]
4.2 Ardeparin versus unfractionated heparin	1	75	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.37 [0.14, 0.99]
4.3 Enoxparin versus unfractionated heparin	1	117	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.34 [0.17, 0.71]
4.4 Certoparin versus unfractionated heparin	1	259	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.63 [0.38, 1.04]
5 Incidence of major haemorrhagic episodes (during initial treatment)	12	6014	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.68 [0.45, 1.03]
5.1 Nadroparin versus unfractionated heparin	4	1436	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.51 [0.18, 1.40]
5.2 Tinzaparin versus unfractionated heparin	2	1044	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.30 [0.12, 0.73]
5.3 Enoxaparin versus unfractionated heparin	3	1034	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.14 [0.50, 2.61]
5.4 Reviparin versus unfractionated heparin	1	1021	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.26 [0.49, 3.19]
5.5 Certoparin versus unfractionated heparin	2	1479	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.62 [0.24, 1.56]
6 Overall mortality at the end of follow-up	14	6984	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.80 [0.65, 0.99]

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 82 thromboembolism (Review)

6.1 Nadroparin versus unfractionated heparin	4	1436	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.76 [0.47, 1.22]
6.2 Tinzaparin versus unfractionated heparin	2	1044	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.63 [0.37, 1.08]
6.3 Ardeparin versus unfractionated heparin	1	80	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.0 [0.0, 0.0]
6.4 Enoxaparin versus unfractionated heparin	4	1934	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.90 [0.63, 1.29]
6.5 Reviparin versus unfractionated heparin	1	1021	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.92 [0.57, 1.47]
6.6 Certoparin versus unfractionated heparin	2	1469	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.70 [0.38, 1.26]

Comparison 9. LMWH versus UFH: all randomised controlled trials that used ISTH definition of major bleeding

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Incidence of major haemorrhagic episodes (during initial treatment)	24	8712	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.71 [0.52, 0.98]
1.1 Nadroparin versus unfractionated heparin	7	1964	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.44 [0.19, 1.01]
1.2 Tinzaparin versus unfractionated heparin	3	1581	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.71 [0.41, 1.22]
1.3 Enoxaparin versus unfractionated heparin	5	1143	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.22 [0.54, 2.75]
1.4 Dalteparin versus unfractionated heparin	4	765	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.15 [0.02, 1.44]
1.5 Reviparin versus unfractionated heparin	1	1021	Peto Odds Ratio (Peto, Fixed, 95% CI)	1.26 [0.49, 3.19]
1.6 Certoparin versus unfractionated heparin	3	2017	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.50 [0.25, 1.00]
1.7 Bemiparin versus unfractionated heparin	1	221	Peto Odds Ratio (Peto, Fixed, 95% CI)	7.39 [0.46, 118.89]

Comparison 10. LMWH versus UFH by year of publication

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Incidence of recurrent venous thromboembolism during initial treatment	18	6238	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.69 [0.49, 0.98]

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 83 thromboembolism (Review)

2 Incidence of recurrent venous thromboembolism at the end	22	9489	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.72 [0.59, 0.88]
of follow-up 3 Incidence of major haemorrhagic episodes (during initial	25	8790	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.69 [0.51, 0.95]
treatment) 4 Overall mortality at the end of follow-up	24	9663	Peto Odds Ratio (Peto, Fixed, 95% CI)	0.84 [0.70, 1.01]

Analysis 1.1. Comparison 1 LMWH versus UFH in patients with venous thromboembolism, Outcome 1 Incidence of recurrent venous thromboembolism during initial treatment.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: I LMWH versus UFH in patients with venous thromboembolism

Outcome: I Incidence of recurrent venous thromboembolism during initial treatment

Study or subgroup	LMWH	UFH	Peto Odds Ratio	Weight	Peto Odds Ratio
,	n/N	n/N	Peto,Fixed,95% Cl	0	Peto,Fixed,95% Cl
I Nadroparin versus unfractior	nated heparin				
Koopman 1996	4/202	5/198	-	7.0 %	0.78 [0.21, 2.92]
Lopaciuk 1992	0/74	1/72	•	0.8 %	0.13 [0.00, 6.64]
Ninet 1991	1/85	2/81		2.3 %	0.48 [0.05, 4.73]
Prandoni 1992	1/85	4/85		3.9 %	0.29 [0.05, 1.72]
Thery 1992	0/35	0/33			Not estimable
Subtotal (95% CI)	481	469	•	14.0 %	0.50 [0.20, 1.26]
Total events: 6 (LMWH), 12 (U	JFH)				
Heterogeneity: $Chi^2 = 1.23$, df	$f = 3 (P = 0.74); I^2 =$	=0.0%			
Test for overall effect: $Z = 1.47$	7 (P = 0.14)				
2 Tinzaparin versus unfractiona	ated heparin				
Simonneau 1997	3/304	2/308	<u> </u>	3.9 %	1.52 [0.26, 8.80]
Subtotal (95% CI)	304	308	-	3.9 %	1.52 [0.26, 8.80]
Total events: 3 (LMWH), 2 (UR	FH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 0.46$	6 (P = 0.64)				
3 Enoxaparin versus unfraction	nated heparin				
Decousus 1998	3/195	8/205		8.5 %	0.41 [0.13, 1.37]
Findik 2002	0/29	1/30	•	0.8 %	0.14 [0.00, 7.06]
Levine 1996	7/247	12/253		14.5 %	0.59 [0.24, 1.48]
			0.001 0.01 0.1 1 10 100 1000		

Favours LMWH Favours UFH

(Continued ...)

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 84 thromboembolism (Review)

Study or subgroup	LMWH n/N	UFH n/N	Peto Odds Ratio Peto,Fixed,95% Cl	Weight	(Continuec Peto Odds Ratio Peto,Fixed,95% Cl
P rez de Llano 2003	3/29	2/21		3.5 %	1.09 [0.17, 6.98]
Simonneau 1993	0/67	2/67		1.6 %	0.13 [0.01, 2.15]
Subtotal (95% CI)	567	576	•	28.9 %	0.51 [0.27, 0.98]
Total events: 13 (LMWH), 25 (3/0		20.9 %	0.91 [0.2/, 0.98]
Heterogeneity: $Chi^2 = 2.18$, df	· /	=0.0%			
Test for overall effect: $Z = 2.03$					
4 Dalteparin versus unfraction	ated heparin				
Fiessinger 1996	1/111	1/120		1.6 %	1.08 [0.07, 17.43]
Lindmarker 1994	0/101	0/103			Not estimable
Meyer 1995	0/29	0/31			Not estimable
Subtotal (95% CI)	241	254	-	1.6 %	1.08 [0.07, 17.43]
Total events: 1 (LMWH), 1 (UI Heterogeneity: not applicable Test for overall effect: Z = 0.06 5 Reviparin versus unfractional Columbus 1997	6 (P = 0.96)	15/511		23.8 %	1.07 [0.52, 2.19]
Subtotal (95% CI) Total events: 16 (LMWH), 15 (Heterogeneity: not applicable Test for overall effect: Z = 0.19 6 Certoparin versus unfractior	9 (P = 0.85)	511		23.8 %	1.07 [0.52, 2.19]
Harenberg 2000a	0/265	3/273		2.4 %	0.14[0.01, 1.34]
Kirchmaier 1998	1/128	3/131		3.1 %	0.37 [0.05, 2.67]
Riess 2003	14/627	15/593	+	22.4 %	0.88 [0.42, 1.84]
Subtotal (95% CI) Total events: 15 (LMWH), 21 (997	•	27.9 %	0.68 [0.35, 1.32]
Heterogeneity: $Chi^2 = 2.72$, df	, ,	=2/%			
Test for overall effect: Z = 1.13 Total (95% CI)	3 (P = 0.26) 3123	3115	•	100.0 %	0.69 [0.49, 0.98]
Total (93.70 CI) Total events: 54 (LMWH), 76 (Heterogeneity: $Chi^2 = 9.77$, df Test for overall effect: $Z = 2.05$	(UFH) F = 14 (P = 0.78); I ²			100.0 /0	0.07 [0.17, 0.76]
Test for subgroup differences: $($	· /	$(P = 0.60), I^2 = 0.0\%$			

0.001 0.01 0.1 1 10 100 1000

Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 85 thromboembolism (Review)

Analysis 1.2. Comparison I LMWH versus UFH in patients with venous thromboembolism, Outcome 2 Incidence of recurrent venous thromboembolism at the end of follow-up.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: I LMWH versus UFH in patients with venous thromboembolism

Outcome: 2 Incidence of recurrent venous thromboembolism at the end of follow-up

Weight	Peto Odds Ratio	UFH	LMWH	Study or subgroup
	Peto,Fixed,95% Cl	n/N	n/N	
			nated heparin	I Nadroparin versus unfraction
4.2 %		13/196	6/98	Belcaro 1999
7.6 %	-	17/198	14/202	Koopman 1996
0.8 %		3/72	0/74	Lopaciuk 1992
4.3 %		12/85	6/85	Prandoni 1992
7.4 %	+	15/360	14/360	Prandoni 2004
24.3 %	•	911	819	Subtotal (95% CI)
			UFH)	Total events: 40 (LMWH), 60 (
		=0.0%	= 4 (P = 0.46); l ² =	Heterogeneity: $Chi^2 = 3.65$, df
			P = 0.14	Test for overall effect: $Z = 1.47$
			ited heparin	2 Tinzaparin versus unfractiona
5.3 %		15/219	6/213	Hull 1992
2.6 %		3/268	7/269	Leizorovicz 2011
2.9 %		6/308	5/304	Simonneau 1997
10.8 %	+	795	786	Subtotal (95% CI)
			UFH)	Total events: 18 (LMWH), 24 (
		-57%	= 2 (P = 0.10); I^2 =	Heterogeneity: $Chi^2 = 4.69$, df
			· /	Test for overall effect: $Z = 0.89$
			ted heparin	3 Ardeparin versus unfractiona
0.3 %	•	1/39	0/41	Goldhaber 1998
0.3 %		39	41	Subtotal (95% CI)
			FH)	Total events: 0 (LMWH), 1 (UF
				Heterogeneity: not applicable
			(P = 0.31)	Test for overall effect: $Z = 1.03$
			ated heparin	4 Enoxaparin versus unfraction
5.5 %	-	12/205	10/195	Decousus 1998
1.0 %		3/30	1/29	Findik 2002
7.5 %	-	17/253	3/247	Levine 1996
7.2 %	+	11/290	21/610	Merli 2001
	4.2 % 7.6 % 0.8 % 4.3 % 7.4 % 24.3 % 5.3 % 2.6 % 2.9 % 10.8 % 0.3 % 0.3 % 0.3 % 0.3 %	Peto,Fixed,95% Cl 42 % 76 % 08 % 43 % 74 % 24.3 % 53 % 26 % 29 % 10.8 % 0.3 % 0.3 % 0.3 % 1.0 % 75 %	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$

Favours LMWH Favours UFH

(Continued ...)

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism (Review)

(... Continued)

Study or subgroup	LMWH	UFH	Peto Odds Ratio	Weight	Peto Odds Ratio
	n/N	n/N	Peto,Fixed,95% Cl		Peto,Fixed,95% Cl
P rez de Llano 2003	0/29	0/21			Not estimable
Simonneau 1993	0/67	3/67		0.8 %	0.13 [0.01, 1.28]
Subtotal (95% CI)	1177	866	•	22.0 %	0.76 [0.49, 1.17]
Total events: 45 (LMWH), 46 (UFH)				
Heterogeneity: $Chi^2 = 3.10$, df	= 4 (P = 0.54); l ² =	=0.0%			
Test for overall effect: $Z = 1.25$. ,				
5 Dalteparin versus unfractiona					
Lindmarker 1994	5/101	3/103		2.1 %	1.71 [0.42, 7.02]
Meyer 1995	0/29	0/31			Not estimable
Subtotal (95% CI)	130	134	-	2.1 %	1.71 [0.42, 7.02]
Total events: 5 (LMWH), 3 (UF	FH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 0.75$	(P = 0.45)				
6 Reviparin versus unfractionat					
Breddin 2001	7/388	24/375	-	7.9 %	0.31 [0.15, 0.63]
Columbus 1997	27/510	25/511	+	13.1 %	1.09 [0.62, 1.90]
Subtotal (95% CI)	898	886	•	21.1 %	0.68 [0.44, 1.05]
Total events: 34 (LMWH), 49 (UFH)				
Heterogeneity: $Chi^2 = 7.38$, df	$= (P = 0.0); ^2 =$	=86%			
Test for overall effect: $Z = 1.74$	(P = 0.082)				
7 Certoparin versus unfraction	ated heparin				
Harenberg 2000a	6/265	15/273		5.4 %	0.42 [0.18, 1.01]
Kirchmaier 1998	2/125	4/124		1.6 %	0.50 [0.10, 2.53]
Riess 2003	22/627	27/593	+	12.5 %	0.76 [0.43, 1.35]
Subtotal (95% CI)	1017	990	•	19.4 %	0.63 [0.40, 0.99]
Total events: 30 (LMWH), 46 (UFH)				
Heterogeneity: $Chi^2 = 1.30$, df	= 2 (P = 0.52); l ² =	=0.0%			
Test for overall effect: $Z = 2.00$	(P = 0.046)				
Total (95% CI)	4868	4621	•	100.0 %	0.72 [0.59, 0.88]
Total events: 172 (LMWH), 22					
Heterogeneity: Chi ² = 22.84, d	,	2 =17%			
Test for overall effect: $Z = 3.23$, ,				
Test for subgroup differences: ($Chi^2 = 2.71, df = 6$	$(P = 0.84), I^2 = 0.0\%$			

0.001 0.01 0.1 1 10 100 1000

Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 87 thromboembolism (Review)

Analysis 1.3. Comparison I LMWH versus UFH in patients with venous thromboembolism, Outcome 3 Incidence of recurrent venous thromboembolism at 1 month follow-up.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: I LMWH versus UFH in patients with venous thromboembolism

Outcome: 3 Incidence of recurrent venous thromboembolism at 1 month follow-up

Study or subgroup	LMWH n/N	UFH n/N	Peto Odds Ratio Peto,Fixed,95% Cl	Weight	Peto Odds Ratio Peto,Fixed,95% Cl
I Nadroparin versus unfractior	nated heparin				
Prandoni 1992	2/85	4/85		8.4 %	0.50 [0.10, 2.55]
Subtotal (95% CI)	85	85	-	8.4 %	0.50 [0.10, 2.55]
Total events: 2 (LMWH), 4 (UF	FH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 0.83$	(P = 0.41)				
2 Enoxaparin versus unfraction	ated heparin				
Levine 1996	7/247	12/253		26.5 %	0.59 [0.24, 1.48]
P rez de Llano 2003	0/29	0/21			Not estimable
Subtotal (95% CI)	276	274	-	26.5 %	0.59 [0.24, 1.48]
Total events: 7 (LMWH), 12 (L	JFH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 1.12$	(P = 0.26)				
3 Reviparin versus unfractionat	ed heparin				
Columbus 1997	25/510	22/511	+	65.0 %	1.15 [0.64, 2.06]
Subtotal (95% CI)	510	511	+	65.0 %	1.15 [0.64, 2.06]
Total events: 25 (LMWH), 22 (UFH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 0.45$	(P = 0.65)				
Total (95% CI)	871	870	+	100.0 %	0.90 [0.56, 1.44]
Total events: 34 (LMWH), 38 (UFH)				
Heterogeneity: $Chi^2 = 1.94$, df	= 2 (P = 0.38); I ² =	=0.0%			
Test for overall effect: $Z = 0.45$	(P = 0.65)				
Test for subgroup differences: ($Chi^2 = 1.94, df = 2$	(P = 0.38), I ² =0.0%			
			0.01 0.1 1 10 100		
			Favours LMWH Favours UFH		

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 88 thromboembolism (Review)

Analysis 1.4. Comparison I LMWH versus UFH in patients with venous thromboembolism, Outcome 4 Incidence of recurrent venous thromboembolism at 3 months' follow-up.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: I LMWH versus UFH in patients with venous thromboembolism

Outcome: 4 Incidence of recurrent venous thromboembolism at 3 months' follow-up

Study or subgroup	LMWH	UFH	Peto Odds Ratio	Weight	Pete Odds Ratio
	n/N	n/N	Peto,Fixed,95% CI		Peto,Fixed,95% (
I Nadroparin versus unfraction	nated heparin				
Belcaro 1999	6/98	13/196		5.9 %	0.92 [0.34, 2.46
Koopman 1996	8/202	10/198		6.5 %	0.78 [0.30, 2.00
Lopaciuk 1992	0/74	3/72		1.1 %	0.13 [0.01, 1.25
Prandoni 1992	4/85	7/85	· _+	3.9 %	0.56 [0.17, 1.89
Prandoni 2004	14/360	15/360		10.4 %	0.93 [0.44, 1.96
Subtotal (95% CI)	819	911	•	27.8 %	0.77 [0.49, 1.21
Total events: 32 (LMWH), 48 (UFH)				• •
Heterogeneity: $Chi^2 = 3.02$, df	= 4 (P = 0.55); l ² =	=0.0%			
Test for overall effect: Z = 1.15	(P = 0.25)				
2 Tinzaparin versus unfractiona	ited heparin				
Hull 1992	6/213	15/219		7.5 %	0.42 [0.17, 1.01
Simonneau 1997	5/304	6/308		4.1 %	0.84 [0.26, 2.77
Subtotal (95% CI)	517	527	•	11.5 %	0.54 [0.26, 1.08
Total events: 11 (LMWH), 21 (UFH)				
Heterogeneity: Chi ² = 0.86, df	= (P = 0.35); ² =	=0.0%			
Test for overall effect: Z = 1.74	(P = 0.083)				
3 Enoxaparin versus unfraction	ated heparin				
Decousus 1998	10/195	12/205		7.8 %	0.87 [0.37, 2.05
Findik 2002	1/29	3/30		1.4 %	0.36 [0.05, 2.70
				1.4 /0	0.50 [0.005; 2.70
Levine 1996	13/247	17/253		1.4 %	0.77 [0.37, 1.62
Levine 1996 Merli 2001	13/247 21/610	7/253 1/290	+		2
			-	10.6 %	0.77 [0.37, 1.62
Merli 2001	21/610	11/290	+ + 	10.6 %	0.77 [0.37, 1.62 0.90 [0.42, 1.92 Not estimab
Merli 2001 P rez de Llano 2003	21/610 0/29	/290 0/2		10.6 %	0.77 [0.37, 1.62 0.90 [0.42, 1.92 Not estimab 0.13 [0.01, 1.28
Merli 2001 P rez de Llano 2003 Simonneau 1993	21/610 0/29 0/67 1177	11/290 0/21 3/67	• • •	10.6 % 10.1 % 1.1 %	0.77 [0.37, 1.62 0.90 [0.42, 1.92
Merli 2001 P rez de Llano 2003 Simonneau 1993 Subtotal (95% CI)	21/610 0/29 0/67 1177 (UFH)	11/290 0/21 3/67 866	•	10.6 % 10.1 % 1.1 %	0.77 [0.37, 1.62 0.90 [0.42, 1.92 Not estimabl 0.13 [0.01, 1.28
Merli 2001 P rez de Llano 2003 Simonneau 1993 Subtotal (95% CI) Total events: 45 (LMWH), 46 (21/610 0/29 0/67 1177 (UFH) = 4 (P = 0.54); I ² =	11/290 0/21 3/67 866		10.6 % 10.1 % 1.1 %	0.77 [0.37, 1.62 0.90 [0.42, 1.92 Not estimab 0.13 [0.01, 1.28
Merli 2001 P rez de Llano 2003 Simonneau 1993 Subtotal (95% CI) Total events: 45 (LMWH), 46 (Heterogeneity: Chi ² = 3.10, df	21/610 0/29 0/67 1177 (UFH) $I = 4 (P = 0.54); I^2 = 0.54$	11/290 0/21 3/67 866		10.6 % 10.1 % 1.1 %	0.77 [0.37, 1.62 0.90 [0.42, 1.92 Not estimab 0.13 [0.01, 1.28

(Continued . . .)

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 89 thromboembolism (Review)

Study or subgroup	LMWH n/N	UFH n/N	Peto Odds Ratio Peto,Fixed,95% Cl	Weight	(Continued) Peto Odds Ratio Peto,Fixed,95% Cl
Meyer 1995	0/29	0/31			Not estimable
Subtotal (95% CI)	29	31			Not estimable
Total events: 0 (LMWH), 0 (UF	Ή)				
Heterogeneity: not applicable					
Test for overall effect: not applie	cable				
5 Reviparin versus unfractionate	ed heparin				
Breddin 2001	7/388	24/375		11.1 %	0.31 [0.15, 0.63]
Columbus 1997	27/510	25/511	+	18.5 %	1.09 [0.62, 1.90]
Subtotal (95% CI)	898	886	•	29.6 %	0.68 [0.44, 1.05]
Total events: 34 (LMWH), 49 (UFH)				
Heterogeneity: Chi ² = 7.38, df	$= (P = 0.0); ^2 =$	=86%			
Test for overall effect: $Z = 1.74$	(P = 0.082)				
Total (95% CI)	3440	3221	•	100.0 %	0.71 [0.56, 0.90]
Total events: 122 (LMWH), 164	4 (UFH)				
Heterogeneity: Chi ² = 15.22, d	f = 13 (P = 0.29); I	2 =15%			
Test for overall effect: $Z = 2.84$	(P = 0.0045)				
Test for subgroup differences: C	$Chi^2 = 0.86, df = 3$	$(P = 0.83), I^2 = 0.0\%$			
			<u> </u>		
			0.01 0.1 1 10 100		
			Favours LMWH Favours UFH		

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 90 thromboembolism (Review)

Analysis 1.5. Comparison I LMWH versus UFH in patients with venous thromboembolism, Outcome 5 Incidence of recurrent venous thromboembolism at 6 months' follow-up.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: I LMWH versus UFH in patients with venous thromboembolism

Outcome: 5 Incidence of recurrent venous thromboembolism at 6 months' follow-up

Study or subgroup	LMWH	UFH	Peto Odds Ratio	Weight	Pete Odds Ratio
	n/N	n/N	Peto,Fixed,95% Cl		Peto,Fixed,95% (
I Nadroparin versus unfraction	ated heparin				
Koopman 1996	14/202	17/198		22.8 %	0.79 [0.38, 1.65
Prandoni 1992	6/85	12/85		12.9 %	0.48 [0.18, 1.26
Subtotal (95% CI)	287	283	•	35.7 %	0.66 [0.37, 1.19
Total events: 20 (LMWH), 29 (,				
Heterogeneity: $Chi^2 = 0.67$, df	, ,	=0.0%			
Test for overall effect: $Z = 1.39$, ,				
2 Dalteparin versus unfractiona					
Lindmarker 1994	5/101	3/103		6.1 %	1.71 [0.42, 7.02
Subtotal (95% CI)	101	103	-	6.1 %	1.71 [0.42, 7.02
Total events: 5 (LMWH), 3 (UF	H)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 0.75$	(P = 0.45)				
3 Certoparin versus unfraction					
Harenberg 2000a	6/265	15/273		16.1 %	0.42 [0.18, 1.01
Kirchmaier 1998	2/125	4/124		4.7 %	0.50 [0.10, 2.53
Riess 2003	22/627	27/593	-	37.4 %	0.76 [0.43, 1.35
Subtotal (95% CI)	1017	990	•	58.2 %	0.63 [0.40, 0.99
Total events: 30 (LMWH), 46 (I	UFH)				
Heterogeneity: $Chi^2 = 1.30$, df	$= 2 (P = 0.52); I^2 =$	=0.0%			
Test for overall effect: $Z = 2.00$	(P = 0.046)				
4 Enoxaparin versus unfraction	ated heparin				
P rez de Llano 2003	0/29	0/31			Not estimab
Subtotal (95% CI)	29	31			Not estimabl
Total events: 0 (LMWH), 0 (UF	H)				
Heterogeneity: not applicable					
Test for overall effect: not applie					
Total (95% CI)	1434	1407	•	100.0 %	0.68 [0.48, 0.96
Total events: 55 (LMWH), 78 (
Heterogeneity: $Chi^2 = 3.76$, df		=0.0%			
Test for overall effect: Z = 2.17 Test for subgroup differences: C	((D - 0.41) + 12 - 0.097			
rest for subgroup amerences: C	∠rii= — 1.78, dī = 2	(r – 0.41), 1 ² –0.0%			

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 91 thromboembolism (Review)

Analysis 1.6. Comparison I LMWH versus UFH in patients with venous thromboembolism, Outcome 6 Reduction in thrombus size (pre- and post-treatment venograms).

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: I LMWH versus UFH in patients with venous thromboembolism

Outcome: 6 Reduction in thrombus size (pre- and post-treatment venograms)

Study or subgroup	LMWH	UFH	Peto Odds Ratio(Non-event)	Weight	Peto Odd Ratio(Non-event
	n/N	n/N	Peto,Fixed,95% Cl	-	Peto,Fixed,95% C
I Nadroparin versus unfractionat	ed heparin				
Lopaciuk 1992	45/68	32/66		5.0 %	0.49 [0.25, 0.96
Ninet 1991	24/78	30/75		5.3 %	1.49 [0.77, 2.89
Prandoni 1992	50/83	36/85	-	6.4 %	0.49 [0.27, 0.90
Thery 1992	29/31	21/21		0.3 %	5.53 [0.32, 95.93
Subtotal (95% CI)	260	247	•	16.9 %	0.72 [0.50, 1.05]
Total events: 148 (LMWH), 119 ((UFH)				
Heterogeneity: Chi ² = 9.45, df =	3 (P = 0.02); I ² =68	3%			
Test for overall effect: $Z = 1.72$ (F	P = 0.086)				
2 Ardeparin versus unfractionated	d heparin				
Goldhaber 1998	31/39	21/36		2.4 %	0.37 [0.14, 0.99
Subtotal (95% CI)	39	36	•	2.4 %	0.37 [0.14, 0.99]
Total events: 31 (LMWH), 21 (UF	FH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 1.97$ (F	P = 0.049)				
3 Enoxaparin versus unfractionate	ed heparin				
Simonneau 1993	35/60	18/57		4.4 %	0.34 [0.17, 0.71
Subtotal (95% CI)	60	57	•	4.4 %	0.34 [0.17, 0.71
Total events: 35 (LMWH), 18 (UF	FH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 2.89$ (F	P = 0.0038)				
4 Dalteparin versus unfractionate	d heparin				
Fiessinger 1996	31/96	41/103		7.0 %	1.38 [0.78, 2.46
Lindmarker 1994	36/91	33/89	+	6.5 %	0.90 [0.49, 1.64
Luomanmaki 1996	47/92	61/98	-	7.1 %	1.57 [0.89, 2.79
Meyer 1995	25/26	26/28		0.4 %	0.54 [0.05, 5.43
	/ 4	11/13		0.6 %	1.47 [0.22, 9.90
Moreno-Palomares 2001					

0.001 0.01 0.1 1 10 100 1000 Favours LMWH Favours UFH

(Continued . . .)

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 92 thromboembolism (Review)

Study or subgroup	LMWH n/N	UFH n/N	Peto Odds Ratio(Non-event) Peto,Fixed,95% Cl	Weight	(Continued) Peto Odds Ratio(Non-event) Peto,Fixed,95% Cl
Total events: 150 (LMWH), 172	(UFH)				
Heterogeneity: $Chi^2 = 2.42$, df =	= 4 (P = 0.66); I ² =0.	0%			
Test for overall effect: $Z = 1.32$ ((P = 0.19)				
5 CY 222 versus unfractionated	heparin				
Faivre 1988	11/30	10/29	+	2.1 %	Peto Odds ight Ratio (Non-event) Peto,Fixed,95% CI 1 % 0.91 [0.32, 2.62] % 0.91 [0.32, 2.62] 4 % 0.59 [0.43, 0.80] 9% 0.59 [0.43, 0.80] 3 % 0.77 [0.49, 1.20] 2 % 0.63 [0.38, 1.04] 9% 0.70 [0.50, 0.98]
Subtotal (95% CI)	30	29	+	2.1 %	0.91 [0.32, 2.62]
Total events: 11 (LMWH), 10 (L	JFH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 0.17$ ((P = 0.86)				
6 Reviparin versus unfractionate	d heparin				
Breddin 2001	175/328	29/32	•	24.4 %	0.59 [0.43, 0.80]
Subtotal (95% CI)	328	321	•	24.4 %	0.59 [0.43, 0.80]
Total events: 175 (LMWH), 129	(UFH)				0.99 [0.13, 0.00]
Heterogeneity: not applicable					
Test for overall effect: $Z = 3.36$ ((P = 0.00078)				
7 Certoparin versus unfractional	ted heparin				
Harenberg 2000a	60/198	48/192	-	11.8 %	0.77 [0.49, 1.20]
Kirchmaier 1998	55/128	42/131	-	9.2 %	0.63 [0.38, 1.04]
Subtotal (95% CI)	326	323	•	21.0 %	0.70 [0.50, 0.98]
Total events: 115 (LMWH), 90 (UFH)				
Heterogeneity: $Chi^2 = 0.34$, df =	= I (P = 0.56); I ² =0.	.0%			
Test for overall effect: $Z = 2.07$ (
8 Bemiparin versus unfractionate	ed heparin				
Kakkar 2003	76/105	51/98	+	7.2 %	0.42 [0.24, 0.74]
Subtotal (95% CI)	105	98	•	7.2 %	0.42 [0.24, 0.74]
Total events: 76 (LMWH), 51 (L	JFH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 2.98$ ((P = 0.0028)				
Total (95% CI)	1467	1442	•	100.0 %	0.71 [0.61, 0.82]
Total events: 741 (LMWH), 610	. ,				
Heterogeneity: $Chi^2 = 33.91$, df		=56%			
Test for overall effect: $Z = 4.45$ ((P < 0.00001)				
Test for subgroup differences: Ch	$hi^2 = 21.70, df = 7$ (1	$P = 0.00$), $I^2 = 68\%$			

0.001 0.01 0.1 1 10 100 1000

Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 93 thromboembolism (Review)

Analysis 1.7. Comparison I LMWH versus UFH in patients with venous thromboembolism, Outcome 7 Incidence of major haemorrhagic episodes (during initial treatment).

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: I LMWH versus UFH in patients with venous thromboembolism

Outcome: 7 Incidence of major haemorrhagic episodes (during initial treatment)

Study or subgroup	LMWH	UFH	Peto Odds Ratio	Weight	Peto Odds Ratio
	n/N	n/N	Peto,Fixed,95% Cl		Peto,Fixed,95% C
I Nadroparin versus unfractio	onated heparin				
Belcaro 1999	0/98	0/196			Not estimable
Koopman 1996	1/202	2/198		1.9 %	0.50 [0.05, 4.85]
Lopaciuk 1992	0/74	1/72	·	0.6 %	0.13 [0.00, 6.64]
Ninet 1991	2/85	4/8		3.8 %	0.48 [0.09, 2.43]
Prandoni 1992	1/85	3/85	<u> </u>	2.5 %	0.36 [0.05, 2.61]
Prandoni 2004	3/360	4/360		4.5 %	0.75 [0.17, 3.32]
Thery 1992	0/35	2/33		1.3 %	0.12 [0.01, 2.02]
Subtotal (95% CI) Total events: 7 (LMWH), 16 (939 (UFH)	1025	•	14.6 %	0.44 [0.19, 1.01]
Heterogeneity: $Chi^2 = 1.71$, c	. ,	=0.0%			
Test for overall effect: $Z = 1.9$	94 (P = 0.052)				
2 Tinzaparin versus unfractior	nated heparin				
Hull 1992	1/213	11/219		7.6 %	0.19 [0.06, 0.59]
Leizorovicz 2011	18/269	15/268	+	20.1 %	1.21 [0.60, 2.44]
Simonneau 1997	3/304	5/308		5.1 %	0.61 [0.15, 2.46]
Subtotal (95% CI)	786	795	•	32.8 %	0.71 [0.41, 1.22]
Total events: 22 (LMWH), 31	(UFH)				
Heterogeneity: $Chi^2 = 7.48$, c	$ff = 2 (P = 0.02); I^2$	=73%			
Test for overall effect: $Z = 1.2$	24 (P = 0.21)				
3 Enoxaparin versus unfractio	nated heparin				
Decousus 1998	7/195	8/205	-	9.4 %	0.92 [0.33, 2.57]
Findik 2002	0/29	0/30			Not estimable
Levine 1996	5/247	3/253		5.1 %	1.70 [0.42, 6.87]
P rez de Llano 2003	1/29	0/21		0.6 %	5.61 [0.11, 297.44]
Simonneau 1993	0/67	0/67			Not estimable
Subtotal (95% CI) Total events: 13 (LMWH), 11	567 (UFH)	576	•	15.1 %	1.22 [0.54, 2.75]
			0.0010.010.11101001000		
			Favours LMWH Favours UFH		

(Continued . . .)

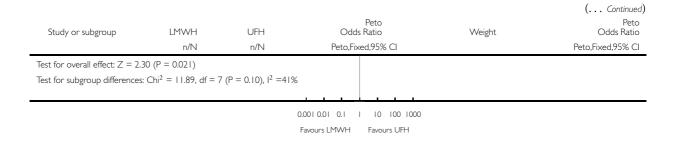
Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 94 thromboembolism (Review)

Study or subgroup	LMWH	UFH	Peto Odds Ratio	Weight	(Continue Peto Odds Ratio
Study or subgroup	n/N	n/N	Peto,Fixed,95% Cl	vveight	Peto,Fixed,95% C
Heterogeneity: Chi ² = 1.08, df					
Test for overall effect: $Z = 0.48$					
4 Dalteparin versus unfractiona	ted heparin				
Fiessinger 1996	0/120	2/133		1.3 %	0.15 [0.01, 2.39
Lindmarker 1994	0/101	0/103			Not estimable
Luomanmaki 1996	0/117	1/131	·	0.6 %	0.15 [0.00, 7.64
Meyer 1995	0/29	0/31			Not estimable
Subtotal (95% CI) Total events: 0 (LMWH), 3 (UF Heterogeneity: Chi ² = 0.00, df	,	398	-	1.9 %	0.15 [0.02, 1.44]
Test for overall effect: $Z = 1.64$, ,				
5 CY 222 versus unfractionated	d heparin				
Faivre 1988	0/33	3/35		1.9 %	0.13 [0.01, 1.34]
Subtotal (95% CI) Total events: 0 (LMWH), 3 (UF	33 ™)	35		1.9 %	0.13 [0.01, 1.34
Heterogeneity: not applicable	(D - 0.000)				
Test for overall effect: Z = 1.71 6 Reviparin versus unfractionate	. ,				
Columbus 1997	10/510	8/511	-	11.5 %	1.26 [0.49, 3.19
Subtotal (95% CI)	510	511	•	11.5 %	1.26 [0.49, 3.19
Total events: 10 (LMWH), 8 (U Heterogeneity: not applicable Test for overall effect: Z = 0.48	,				
7 Certoparin versus unfraction	. ,				
Harenberg 2000a	4/265	11/273		9.4 %	0.40 [0.14, 1.10
Kirchmaier 1998	1/128	4/131		3.2 %	0.30 [0.05, 1.77
Riess 2003	6/627	7/593	+	8.3 %	0.81 [0.27, 2.41
Subtotal (95% CI) Total events: 11 (LMWH), 22 (1020 UFH)	997	•	21.0 %	0.50 [0.25, 1.00
Heterogeneity: $Chi^2 = 1.26$, df Test for overall effect: $Z = 1.95$	(P = 0.052)	=0.0%			
8 Bemiparin versus unfractiona Kakkar 2003	ted heparin 2/111	0/110		1.3 %	7.39 [0.46, 118.89
Subtotal (95% CI)	111	110		1.3 %	7.39 [0.46, 118.89
Total events: 2 (LMWH), 0 (UF Heterogeneity: not applicable	,				
Test for overall effect: Z = 1.41 Total (95% CI)	(P = 0.16) 4333	4447	•	100.0 %	0.69 [0.50, 0.95
Total events: 65 (LMWH), 94 (Heterogeneity: Chi ² = 23.42, d	UFH)				

Favours LMWH Favours UFH

(Continued . . .)

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 95 thromboembolism (Review)



Analysis I.8. Comparison I LMWH versus UFH in patients with venous thromboembolism, Outcome 8 Overall mortality at the end of follow-up.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism Comparison: I LMWH versus UFH in patients with venous thromboembolism

Outcome: 8 Overall mortality at the end of follow-up

Study or subgroup	LMWH	UFH	Peto Odds Ratio	Weight	Peto Odds Ratio
	n/N	n/N	Peto,Fixed,95% CI		Peto,Fixed,95% CI
I Nadroparin versus unfraction	nated heparin				
Koopman 1996	14/202	16/198	-	6.1 %	0.85 [0.40, 1.78]
Lopaciuk 1992	0/74	1/72	·	0.2 %	0.13 [0.00, 6.64]
Prandoni 1992	6/85	12/85		3.6 %	0.48 [0.18, 1.26]
Prandoni 2004	12/360	12/360	+	5.1 %	1.00 [0.44, 2.26]
Thery 1992	1/35	1/33		0.4 %	0.94 [0.06, 15.40]
Subtotal (95% CI)	756	748	•	15.4 %	0.77 [0.48, 1.22]
Total events: 33 (LMWH), 42	(UFH)				
Heterogeneity: Chi ² = 2.19, df	$f = 4 (P = 0.70); I^2 =$	=0.0%			
Test for overall effect: $Z = 1.12$	2 (P = 0.26)				
2 Tinzaparin versus unfraction	ated heparin				
Hull 1992	10/213	21/219	-	6.3 %	0.48 [0.23, 1.00]
Leizorovicz 2011	31/269	17/268		9.6 %	1.89 [1.04, 3.41]
Simonneau 1997	12/304	4/308	-	5.5 %	0.86 [0.39, 1.89]
Subtotal (95% CI)	786	795	+	21.4 %	1.03 [0.69, 1.53]

0.001 0.01 0.1 1 10 100 1000

Favours LMWH Favours UFH

(Continued . . .)

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 96 thromboembolism (Review)

Study or subgroup	LMWH	UFH	Peto Odds Ratio	Weight	(Continue Peto Odds Ratio
	n/N	n/N	Peto,Fixed,95% Cl		Peto,Fixed,95% C
Total events: 53 (LMWH), 52 (Heterogeneity: $Chi^2 = 8.40$, df Test for overall effect: $Z = 0.15$ 3 Ardeparin versus unfractiona	$= 2 (P = 0.01); I^2 = (P = 0.88)$	=76%			
Goldhaber 1998	0/41	0/39			Not estimable
Subtotal (95% CI)	41	39			Not estimable
Total events: 0 (LMWH), 0 (UF Heterogeneity: not applicable Test for overall effect: not appli 4 Enoxaparin versus unfraction:	^E H) cable				
Decousus 1998	40/195	43/205	+	14.5 %	0.97 [0.60, 1.58
Findik 2002	0/29	0/30			Not estimable
Levine 1996	11/247	17/253	-	5.8 %	0.65 [0.30, 1.40
Merli 2001	18/610	9/290	+	5.0 %	0.95 [0.42, 2.15
P rez de Llano 2003	1/29	0/21		0.2 %	5.61 [0.11, 297.44
Simonneau 1993	3/67	2/67		1.1 %	1.51 [0.25, 8.96
Subtotal (95% CI)	1177	866	•	26.6 %	0.92 [0.64, 1.31
Test for overall effect: Z = 0.49 5 Dalteparin versus unfractiona Lindmarker 1994	, ,	3/103		1.1 %	0.68 [0.12, 3.99
Luomanmaki 1996	1/110	4/116		1.1 %	0.31 [0.05, 1.82
Meyer 1995	0/29	0/31			Not estimabl
Subtotal (95% CI)	240	250	•	2.2 %	0.46 [0.13, 1.60
Total events: 3 (LMWH), 7 (UF Heterogeneity: Chi ² = 0.37, df Test for overall effect: Z = 1.22 6 Reviparin versus unfractionate Breddin 2001	= I (P = 0.54); I ² = (P = 0.22)	=0.0% 1/375		4.3 %	0.79 [0.32, 1.91
Columbus 1997	36/510	39/511	+	15.3 %	0.92 [0.57, 1.47
Subtotal (95% CI)	898	886	•	19.6 %	0.89 [0.59, 1.35
Total events: 45 (LMWH), 50 (I Heterogeneity: $Chi^2 = 0.09$, df Test for overall effect: $Z = 0.56$ 7 Certoparin versus unfraction:	UFH) = I (P = 0.76); I ² = (P = 0.58)				
Harenberg 2000a	6/265	15/273		4.4 %	0.42 [0.18, 1.01
Kirchmaier 1998	8/125	10/124	<u> </u>	3.7 %	0.78 [0.30, 2.03
			0.001 0.01 0.1 1 10 100 1000 Favours LMWH Favours UFH		

(Continued . . .)

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 97 thromboembolism (Review)

Study or subgroup	LMWH	UFH	Peto Odds Ratio	Weight	(Continued) Peto Odds Ratio
D: 2002	n/N	n/N	Peto,Fixed,95% Cl	F 0.0/	Peto,Fixed,95% Cl
Riess 2003	11/627	16/593		5.8 %	0.65 [0.30, 1.39]
Subtotal (95% CI)	1017	990	•	13.9 %	0.59 [0.36, 0.97]
Total events: 25 (LMWH), 41 (UFH)				
Heterogeneity: Chi ² = 0.94, df	= 2 (P = 0.62); I ² =	0.0%			
Test for overall effect: $Z = 2.07$	r (P = 0.038)				
8 Bemiparin versus unfractiona	ted heparin				
Kakkar 2003	2/89	2/85		0.9 %	0.95 [0.13, 6.90]
Subtotal (95% CI)	89	85	-	0.9 %	0.95 [0.13, 6.90]
Total events: 2 (LMWH), 2 (UF	FH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 0.05$	(P = 0.96)				
Total (95% CI)	5004	4659	•	100.0 %	0.84 [0.70, 1.01]
Total events: 234 (LMWH), 26	5 (UFH)				
Heterogeneity: $Chi^2 = 18.23$, c	$If = 20 (P = 0.57); I^2$	=0.0%			
Test for overall effect: $Z = 1.82$	(P = 0.068)				
Test for subgroup differences: ($Chi^2 = 4.29, df = 6$	$P = 0.64$), $I^2 = 0.0\%$			
			0.00 0.0 0.1 0 00 000		

Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 98 thromboembolism (Review)

Analysis 2.1. Comparison 2 LMWH versus UFH in patients with proximal deep venous thrombosis, Outcome I Incidence of recurrent venous thromboembolism at the end of follow-up.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 2 LMWH versus UFH in patients with proximal deep venous thrombosis

Outcome: I Incidence of recurrent venous thromboembolism at the end of follow-up

Study or subgroup	LMWH	UFH	Peto Odds Ratio	Weight	Petc Odds Ratio
	n/N	n/N	Peto,Fixed,95% Cl		Peto,Fixed,95% C
I Nadroparin versus unfraction	nated heparin				
Belcaro 1999	6/98	13/196		7.6 %	0.92 [0.34, 2.46
Koopman 1996	14/202	17/198		13.7 %	0.79 [0.38, 1.65
Prandoni 1992	6/85	12/85		7.7 %	0.48 [0.18, 1.26
Subtotal (95% CI)	385	479	•	29.1 %	0.72 [0.44, 1.19]
Total events: 26 (LMWH), 42 ((UFH)				
Heterogeneity: Chi ² = 0.99, df	= 2 (P = 0.61); I ² =	=0.0%			
Test for overall effect: $Z = 1.28$	8 (P = 0.20)				
2 Tinzaparin versus unfractiona	ated heparin				
Hull 1992	6/213	15/219		9.6 %	0.42 [0.17, 1.01]
Subtotal (95% CI)	213	219	•	9.6 %	0.42 [0.17, 1.01]
Total events: 6 (LMWH), 15 (L	JFH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 1.95$	6 (P = 0.052)				
3 Enoxaparin versus unfraction	lated heparin				
Levine 1996	13/247	17/253		13.5 %	0.77 [0.37, 1.62
Simonneau 1993	0/67	3/67		1.4 %	0.13 [0.01, 1.28
Subtotal (95% CI)	314	320	•	14.9 %	0.65 [0.32, 1.32
Total events: 13 (LMWH), 20 ((UFH)				
Heterogeneity: $Chi^2 = 2.10$, df	$= (P = 0. 5); ^2 =$	=52%			
Test for overall effect: $Z = 1.19$	P (P = 0.23)				
4 Reviparin versus unfractionat	ed heparin				
Breddin 2001	7/388	24/375		14.2 %	0.31 [0.15, 0.63
Subtotal (95% CI)	388	375	•	14.2 %	0.31 [0.15, 0.63]
Total events: 7 (LMWH), 24 (L	JFH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 3.21$	(P = 0.0013)				
5 Certoparin versus unfraction					
Harenberg 2000a	6/265	15/273		9.7 %	0.42 [0.18, 1.01
Riess 2003	22/627	27/593	-	22.5 %	0.76 [0.43, 1.35
Subtotal (95% CI)	892	866	•	32.2 %	0.64 [0.40, 1.03
			0.01 0.1 1 10 100		
			Favours LMWH Favours UFH		
					/

(Continued \dots)

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 99 thromboembolism (Review)

Study or subgroup	LMWH	UFH	Peto Odds Ratio	Weight	(<i>Continued</i>) Peto Odds Ratio
	n/N	n/N	Peto,Fixed,95% Cl		Peto,Fixed,95% Cl
Total events: 28 (LMWH), 42 ((UFH)				
Heterogeneity: Chi ² = 1.23, df	$= 1 (P = 0.27); I^2 =$	=18%			
Test for overall effect: $Z = 1.84$	+ (P = 0.066)				
6 Bemiparin versus unfractiona	ited heparin				
Kakkar 2003	0/111	0/110			Not estimable
Subtotal (95% CI)	111	110			Not estimable
Total events: 0 (LMWH), 0 (UI	FH)				
Heterogeneity: not applicable					
Test for overall effect: not appli	icable				
Total (95% CI)	2303	2369	•	100.0 %	0.57 [0.44, 0.75]
Total events: 80 (LMWH), 143	(UFH)				
Heterogeneity: $Chi^2 = 8.80$, df	= 8 (P = 0.36); I ² =	-9%			
Test for overall effect: $Z = 4.01$	(P = 0.000062)				
Test for subgroup differences: ($Chi^2 = 4.48, df = 4$	$(P = 0.34), ^2 = \%$			

0.01 0.1 1 10 100 Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 100 thromboembolism (Review)

Analysis 2.2. Comparison 2 LMWH versus UFH in patients with proximal deep venous thrombosis, Outcome 2 Incidence of recurrent deep venous thrombosis at the end of follow-up.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 2 LMWH versus UFH in patients with proximal deep venous thrombosis

Outcome: 2 Incidence of recurrent deep venous thrombosis at the end of follow-up

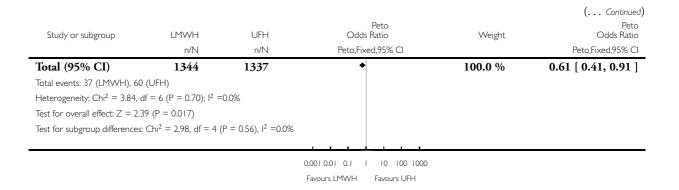
Pe [.] Odds Rat	Weight	Peto Odds Ratio	UFH	LMWH	Study or subgroup
Peto,Fixed,95%		Peto,Fixed,95% Cl	n/N	n/N	
				ated heparin	I Nadroparin versus unfractiona
0.99 [0.31, 3.17	12.2 %	+	6/97	6/98	Belcaro 1999
0.81 [0.34, 1.91	22.4 %	-	12/198	10/202	Koopman 1996
0.41 [0.09, 1.86	7.3 %		5/85	2/85	Prandoni 1992
0.76 [0.41, 1.43	41.9 %	•	380	385	Subtotal (95% CI)
			0.0%	= 2 (P = 0.65); I ² = (P = 0.40)	Total events: 18 (LMWH), 23 (L Heterogeneity: $Chi^2 = 0.85$, df = Test for overall effect: Z = 0.85
0.37 [0.12, 1.16	12.6 %		9/219	ed neparin 3/213	2 Tinzaparin versus unfractionat Hull 1992
0.37 [0.12, 1.16]	12.6 %	-	219	213	Subtotal (95% CI) Total events: 3 (LMWH), 9 (UFI
				(P = 0.088)	Heterogeneity: not applicable Test for overall effect: Z = 1.71 3 Enoxaparin versus unfractiona
0.74 [0.34, 1.63	26.6 %	-	15/253	/247	Levine 1996
0.74 [0.34, 1.63	26.6 %	•	253	(P = 0.46)	Subtotal (95% CI) Total events: 11 (LMWH), 15 (L Heterogeneity: not applicable Test for overall effect: Z = 0.74 4 Reviparin versus unfractionate
0.45 [0.17, 1.21	16.9 %		11/375	5/388	Breddin 2001
0.45 [0.17, 1.21	16.9 %	•	375	(P = 0.11)	Subtotal (95% CI) Total events: 5 (LMWH), 11 (UI Heterogeneity: not applicable Test for overall effect: Z = 1.58 5 Bemiparin versus unfractionat
0.13 [0.01, 2.14	2.1 %		2/110	0/111	Kakkar 2003
0.13 [0.01, 2.14	2.1 %		110	,	Subtotal (95% CI) Total events: 0 (LMWH), 2 (UFI Heterogeneity: not applicable Test for overall effect: Z = 1.42

(Continued ...)

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 101 thromboembolism (Review)

Favours LMWH Favours UFH

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Analysis 2.3. Comparison 2 LMWH versus UFH in patients with proximal deep venous thrombosis, Outcome 3 Incidence of pulmonary embolism at the end of follow-up.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 2 LMWH versus UFH in patients with proximal deep venous thrombosis

Outcome: 3 Incidence of pulmonary embolism at the end of follow-up

Study or subgroup	LMWH	UFH	Peto Odds Ratio	Weight	Peto Odds Ratio
otady of stopping	n/N	n/N	Peto,Fixed,95% Cl	, togic	Peto,Fixed,95% Cl
I Nadroparin versus unfractio	nated heparin				
Koopman 1996	4/202	5/198		13.9 %	0.78 [0.21, 2.92]
Prandoni 1992	4/85	7/85		16.3 %	0.56 [0.17, 1.89]
Subtotal (95% CI)	287	283	•	30.3 %	2
Total events: 8 (LMWH), 12 (U	JFH)				
Heterogeneity: $Chi^2 = 0.13$, d	$f = (P = 0.72); ^2 =$	=0.0%			
Test for overall effect: $Z = 0.92$	3 (P = 0.35)				
2 Tinzaparin versus unfraction	ated heparin				
Hull 1992	3/213	6/219		14.0 %	0.52 [0.14, 1.95]
Subtotal (95% CI)	213	219	•	14.0 %	0.52 [0.14, 1.95]
Total events: 3 (LMWH), 6 (U	FH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 0.97$	7 (P = 0.33)				
3 Enoxaparin versus unfractior	nated heparin				
			0.001 0.01 0.1 1 10 100 1000		
			Favours LMWH Favours UFH		

(Continued . . .)

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 102 thromboembolism (Review)

(... Continued)

(Continue Petc Odds Ratic Peto,Fixed,95% Cl	Weight	Peto Odds Ratio Peto,Fixed,95% Cl	UFH n/N	LMWH n/N	Study or subgroup
0.52 [0.05, 5.07]	4.7 %		2/253	1/247	Levine 1996
0.52 [0.05, 5.07]	4.7 %	-	253	247	Subtotal (95% CI)
				H)	Total events: I (LMWH), 2 (UF Heterogeneity: not applicable
				· /	Test for overall effect: $Z = 0.56$
		_		·	4 Reviparin versus unfractionate
0.27 [0.10, 0.73]	24.8 %		13/375	3/388	Breddin 2001
0.27 [0.10, 0.73]	24.8 %	•	375	388	Subtotal (95% CI)
				IFH)	Total events: 3 (LMWH), 13 (U
					Heterogeneity: not applicable
				(P = 0.0095)	Test for overall effect: $Z = 2.59$
					ated heparin
0.32 [0.11, 0.92]	21.6 %		11/273	3/265	Harenberg 2000a
0.32 [0.11, 0.92]	21.6 %	•	273	265	Subtotal (95% CI)
				IFH)	Total events: 3 (LMWH), 11 (U
					Heterogeneity: not applicable
				(P = 0.035)	Test for overall effect: $Z = 2.11$
				ted heparin	6 Bemiparin versus unfractionat
1.94 [0.20, 18.86]	4.7 %		1/110	2/111	Kakkar 2003
1.94 [0.20, 18.86]	4.7 %	-	110	111	Subtotal (95% CI)
				H)	Total events: 2 (LMWH), 1 (UF
					Heterogeneity: not applicable
				(P = 0.57)	Test for overall effect: Z = 0.57
0.45 [0.28, 0.74]	100.0 %	•	1513	1511	Total (95% CI)
				UFH)	Total events: 20 (LMWH), 45 (I
			=0.0%	$= 6 (P = 0.69); I^2 =$	Heterogeneity: $Chi^2 = 3.87$, df
				` '	Test for overall effect: $Z = 3.14$
			$(P = 0.59), I^2 = 0.0\%$	$Chi^2 = 3.74, df = 5$	Test for subgroup differences: C
		<u> </u>			

Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 103 thromboembolism (Review)

Analysis 2.4. Comparison 2 LMWH versus UFH in patients with proximal deep venous thrombosis, Outcome 4 Reduction in thrombus size (pre- and post-treatment venograms).

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 2 LMWH versus UFH in patients with proximal deep venous thrombosis

Outcome: 4 Reduction in thrombus size (pre- and post-treatment venograms)

Study or subgroup	LMWH	UFH	Peto Odds Ratio(Non-event)	Weight	Peto Odds Ratio(Non-event)
	n/N	n/N	Peto,Fixed,95% Cl	-	Peto,Fixed,95% Cl
I Bemiparin versus unfractionated	d heparin				
Kakkar 2003	76/105	51/98		91.9 %	0.42 [0.24, 0.74]
Subtotal (95% CI)	105	98	•	91.9 %	0.42 [0.24, 0.74]
Total events: 76 (LMWH), 51 (UF	FH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 2.98$ (F	P = 0.0028)				
2 Dalteparin versus unfractionate	d heparin				
Moreno-Palomares 2001	/ 4	/ 3		8.1 %	1.47 [0.22, 9.90]
Subtotal (95% CI)	14	13	-	8.1 %	1.47 [0.22, 9.90]
Total events: (LMWH), (UF	FH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 0.40$ (F	P = 0.69)				
Total (95% CI)	119	111	•	100.0 %	0.47 [0.27, 0.80]
Total events: 87 (LMWH), 62 (UF	FH)				
Heterogeneity: $Chi^2 = 1.52$, df =	$ (P = 0.22); ^2 = 34$	4%			
Test for overall effect: $Z = 2.75$ (F	P = 0.0060)				
Test for subgroup differences: Chi	² = 1.52, df = 1 (P	= 0.22), l ² =34%			

0.002 0.1 I 10 500 Favours LMWH Favours UFH

Analysis 2.5. Comparison 2 LMWH versus UFH in patients with proximal deep venous thrombosis, Outcome 5 Incidence of major haemorrhagic episodes (during initial treatment).

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 2 LMWH versus UFH in patients with proximal deep venous thrombosis

Outcome: 5 Incidence of major haemorrhagic episodes (during initial treatment)

Study or subgroup	LWMH	UFH	Peto Odds Ratio	Weight	Pet Odds Rati
	n/N	n/N	Peto,Fixed,95% Cl		Peto,Fixed,95% (
I Nadroparin versus unfractional	ted heparin				
Belcaro 1999	0/98	0/97			Not estimab
Koopman 1996	1/202	2/198		5.5 %	0.50 [0.05, 4.85
Prandoni 1992	1/85	3/85		7.3 %	0.36 [0.05, 2.61
Subtotal (95% CI)	385	380	-	12.8 %	0.42 [0.09, 1.85
Total events: 2 (LWMH), 5 (UFH Heterogeneity: $Chi^2 = 0.05$, df = Test for overall effect: Z = 1.15 (P = 0.25)	=0.0%			
2 Tinzaparin versus unfractionate Hull 1992	ed heparin 1/213	11/219		21.7 %	0.19 [0.06, 0.59
Subtotal (95% CI)	213	219	•	21.7 %	0.19 [0.06, 0.59
Total events: I (LWMH), II (UFI Heterogeneity: not applicable Test for overall effect: Z = 2.88 (3 Enoxaparin versus unfractionat	P = 0.0040)				
Levine 1996	5/247	3/253		14.6 %	1.70 [0.42, 6.87
Simonneau 1993	0/67	0/67			Not estimab
Subtotal (95% CI) Total events: 5 (LWMH), 3 (UFH Heterogeneity: not applicable Test for overall effect: Z = 0.75 (4 Certoparin versus unfractionat	P = 0.46)	320	-	14.6 %	1.70 [0.42, 6.87
Harenberg 2000a	4/265	11/273		27.1 %	0.40 [0.14, 1.10
Riess 2003	6/627	7/593	-	23.8 %	0.81 [0.27, 2.41
Subtotal (95% CI)	892	866	•	50.9 %	0.55 [0.26, 1.17
Total events: 10 (LWMH), 18 (UI Heterogeneity: $Chi^2 = 0.88$, df = Test for overall effect: Z = 1.55 (: (P = 0.35); ² =	=0.0%			
Total (95% CI) Total events: 18 (LWMH), 37 (UI Heterogeneity: Chi ² = 6.87, df =	1804 FH) 5 (P = 0.23); I ² =	1785	•	100.0 %	0.50 [0.29, 0.85
Test for overall effect: Z = 2.57 (Test for subgroup differences: Ch	,	$(P = 0.11), 1^2 = 50\%$			

Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 105 thromboembolism (Review)

Analysis 2.6. Comparison 2 LMWH versus UFH in patients with proximal deep venous thrombosis, Outcome 6 Overall mortality at the end of follow-up.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism Comparison: 2 LMWH versus UFH in patients with proximal deep venous thrombosis

Outcome: 6 Overall mortality at the end of follow-up

0.48 [0.18, 1.26	16.0 %	Peto,Fixed,95% Cl	n/N	n/N	
0.48 [0.18, 1.26					
0.48 [0.18, 1.26				ated heparin	I Nadroparin versus unfraction
L .	9.3 %	1	16/198	14/202	Koopman 1996
0.69 [0.38, 1.24			12/85	6/85	Prandoni 1992
	25.3 %	•	283	287	Subtotal (95% CI)
				JFH)	Total events: 20 (LWMH), 28 (U
			=0.0%		Heterogeneity: $Chi^2 = 0.85$, df
				(P = 0.21)	Test for overall effect: Z = 1.25
				ted heparin	2 Tinzaparin versus unfractionat
0.48 [0.23, 1.00	16.6 %		21/219	10/213	Hull 1992
0.48 [0.23, 1.00]	16.6 %	•	219	213	Subtotal (95% CI)
				JFH)	Total events: 10 (LWMH), 21 (U
					Heterogeneity: not applicable
				(P = 0.049)	Test for overall effect: Z = 1.97
				ated heparin	3 Enoxaparin versus unfractiona
0.65 [0.30, 1.40	15.2 %		17/253	11/247	Levine 1996
1.51 [0.25, 8.96	2.8 %		2/67	3/67	Simonneau 1993
0.74 [0.37, 1.50]	18.0 %	•	320	314	Subtotal (95% CI)
				JFH)	Total events: 14 (LWMH), 19 (U
			=0.0%	$= (P = 0.40); ^2 =$	Heterogeneity: $Chi^2 = 0.72$, df
				(P = 0.40)	Test for overall effect: Z = 0.83
				ed heparin	4 Reviparin versus unfractionate
0.79 [0.32, 1.91	11.2 %		/375	9/388	Breddin 2001
0.79 [0.32, 1.91	11.2 %	•	375	388	Subtotal (95% CI)
				FH)	Total events: 9 (LWMH), 11 (U
					Heterogeneity: not applicable
				(P = 0.60)	Test for overall effect: $Z = 0.53$
				ated heparin	5 Certoparin versus unfractiona
0.42 [0.18, 1.01	11.6 %		15/273	6/265	Harenberg 2000a
0.65 [0.30, 1.39	15.2 %		16/593	11/627	Riess 2003

(Continued . . .)

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 106 thromboembolism (Review)

Study or subgroup	LWMH	UFH	Peto Odds Ratio	Weight	(Continued) Peto Odds Ratio
	n/N	n/N	Peto,Fixed,95% Cl		Peto,Fixed,95% Cl
Subtotal (95% CI)	892	866	•	26.8 %	0.54 [0.30, 0.96]
Total events: 17 (LWMH), 31 (I	UFH)				
Heterogeneity: $Chi^2 = 0.5 I$, df	$= (P = 0.47); ^2 =$	=0.0%			
Test for overall effect: $Z = 2.11$	(P = 0.034)				
6 Bemiparin versus unfractiona	ted heparin				
Kakkar 2003	2/89	2/85		2.3 %	0.95 [0.13, 6.90]
Subtotal (95% CI)	89	85	-	2.3 %	0.95 [0.13, 6.90]
Total events: 2 (LWMH), 2 (UF	Ή)				
Heterogeneity: not applicable					
Test for overall effect: Z = 0.05	(P = 0.96)				
Total (95% CI)	2183	2148	•	100.0 %	0.63 [0.47, 0.85]
Total events: 72 (LWMH), 112	(UFH)				
Heterogeneity: $Chi^2 = 3.60$, df	= 8 (P = 0.89); l ² =	=0.0%			
Test for overall effect: $Z = 3.06$	(P = 0.0022)				
Test for subgroup differences: C	$Chi^2 = 1.52, df = 5$	$(P = 0.9 I), I^2 = 0.0\%$			
			0.01 0.1 1 10 10	0	
			Favours LWMH Favours UFH		

Analysis 3.1. Comparison 3 LMWH versus UFH in patients with pulmonary embolism, Outcome I Incidence of recurrent venous thromboembolism at the end of follow-up.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 3 LMWH versus UFH in patients with pulmonary embolism

Outcome: I Incidence of recurrent venous thromboembolism at the end of follow-up

LMWH	-	UFH	Peto Odds Ratio	Weight	Odds Ratic
n/N	N	n/N	Peto,Fixed,95% Cl		Peto,Fixed,95% C
d heparin					
5/304	4	6/308		23.7 %	0.84 [0.26, 2.77]
0/35	5	0/33			Not estimable
339	9	341	•	23.7 %	0.84 [0.26, 2.77]
)					
P = 0.78)					
ed heparin					
1/29	9	3/30		8.3 %	0.36 [0.05, 2.70]
10/199	9	4/88	-	24.9 %	1.11 [0.35, 3.55]
3/29	9	2/21		9.8 %	1.09 [0.17, 6.98]
257	7	139	+	43.1 %	0.89 [0.37, 2.16]
H)					
2 (P = 0.62	2); I ² =0.0%	Ś			
2 (P = 0.62 P = 0.80)	52); I ² =0.0%	ó			
	2); I ² =0.0%	5			
P = 0.80)	,	0/31			Not estimable
P = 0.80) d heparin	9				Not estimable Not estimable
P = 0.80) d heparin 0/29	9	0/31			
P = 0.80) d heparin 0/29 29	9	0/31			
P = 0.80) d heparin 0/29 29	9	0/31			
P = 0.80) d heparin 0/29 29	9	0/31			
P = 0.80) d heparin 0/29 29) ble	9 9	0/31	-	33.2 %	
P = 0.80) d heparin 0/29 29) ble heparin	9 9 8	0/31 31	+	33.2 % 33.2 %	Not estimable
 P = 0.80) d heparin 0/29 29 ble heparin 8/138 	9 9 8	0/31 31 8/133	•		Not estimable 0.96 [0.35, 2.64]
2 = 0.80) d heparin 0/29 29) ble heparin 8/138	9 9 8	0/31 31 8/133	•		Not estimable 0.96 [0.35, 2.64]
2 = 0.80) d heparin 0/29 29) ble heparin 8/138	9 9 8	0/31 31 8/133	-		Not estimable 0.96 [0.35, 2.64]
2 = 0.80) d heparin 0/29) ble heparin 8/138 138	9 9 8 8	0/31 31 8/133	+		Not estimable 0.96 [0.35, 2.64 0.96 [0.35, 2.64]
2 = 0.80) d heparin 0/29) ble heparin 8/138) 2 = 0.94)	9 9 8 8	0/31 31 8/133 133	•	33.2 %	Not estimable 0.96 [0.35, 2.64 0.96 [0.35, 2.64]
P = 0.80) d heparin 0/25 29) ble heparin 8/138 138) P = 0.94) 763 TH)	9 9 8 8	0/31 31 8/133 133 644	•	33.2 %	Not estimable 0.96 [0.35, 2.64 0.96 [0.35, 2.64]
P = 0.80) d heparin 0/25 29) ble heparin 8/138 138) P = 0.94) 763 TH)	9 9 8 8 8	0/31 31 8/133 133 644	•	33.2 %	Not estimable 0.96 [0.35, 2.64]

Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 108 thromboembolism (Review)

Analysis 3.2. Comparison 3 LMWH versus UFH in patients with pulmonary embolism, Outcome 2 Reduction in thrombus size (pre- and post-treatment venograms).

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 3 LMWH versus UFH in patients with pulmonary embolism

Outcome: 2 Reduction in thrombus size (pre- and post-treatment venograms)

Study or subgroup	LMWH	UFH	Peto Odds Ratio(Non-event)	Weight	Peto Odds Ratio(Non-event)
, 5 1	n/N	n/N	Peto,Fixed,95% Cl		Peto,Fixed,95% Cl
I Nadroparin versus unfractio	nated heparin				
Thery 1992	29/31	21/21		39.6 %	5.53 [0.32, 95.93]
Subtotal (95% CI)	31	21		39.6 %	5.53 [0.32, 95.93]
Total events: 29 (LMWH), 21	(UFH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 1.1$	8 (P = 0.24)				
2 Dalteparin versus unfraction	ated heparin				
Meyer 1995	25/26	26/28		60.4 %	0.54 [0.05, 5.43]
Subtotal (95% CI)	26	28	-	60.4 %	0.54 [0.05, 5.43]
Total events: 25 (LMWH), 26	(UFH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 0.5$	2 (P = 0.60)				
Total (95% CI)	57	49	*	100.0 %	1.36 [0.23, 8.16]
Total events: 54 (LMWH), 47	(UFH)				
Heterogeneity: Chi ² = 1.55, d	$f = (P = 0.2); ^2 =$	=35%			
Test for overall effect: $Z = 0.3$	3 (P = 0.74)				
Test for subgroup differences:	$Chi^2 = 1.55, df = 1$	$(P = 0.21), I^2 = 35\%$			

0.001 0.01 0.1 1 10 100 1000 Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 109 thromboembolism (Review)

Analysis 3.3. Comparison 3 LMWH versus UFH in patients with pulmonary embolism, Outcome 3 Mean change in pulmonary vascular obstruction severity score.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 3 LMWH versus UFH in patients with pulmonary embolism

Outcome: 3 Mean change in pulmonary vascular obstruction severity score

Study or subgroup	LMWH N	Mean(SD)	UFH N	Mean(SD)	Mean Difference IV,Fixed,95% Cl	Weight	Mean Difference IV,Fixed,95% CI
I Nadroparin versus unfract	ionated hep	parin					
Thery 1992	31	20.48 (2.28)	21	23.76 (2.29)		96.8 %	-3.28 [-4.55, -2.01]
Subtotal (95% CI)	31		21		•	96.8 %	-3.28 [-4.55, -2.01]
Heterogeneity: not applicabl	e						
Test for overall effect: $Z = 5$.	.08 (P < 0.0	0001)					
2 Dalteparin versus unfractio	onated hepa	ırin					
Meyer 1995	26	17 (13)	28	16 (13)	+	3.2 %	1.00 [-5.94, 7.94]
Subtotal (95% CI)	26		28		+	3.2 %	1.00 [-5.94, 7.94]
Heterogeneity: not applicabl	e						
Test for overall effect: $Z = 0$.	.28 (P = 0.7	(8)					
Total (95% CI)	57		49		•	100.0 %	-3.14 [-4.39, -1.90]
Heterogeneity: Chi ² = 1.41,	df = 1 (P =	: 0.23); l ² =29%					
Test for overall effect: $Z = 4$.	.94 (P < 0.0	0001)					
Test for subgroup difference:	s: Chi ² = 1.4	41, df = 1 (P = 0.	23), I ² =2	9%			

-100 -50 0 50 100

Favours LMWH Favours UFH

Analysis 3.4. Comparison 3 LMWH versus UFH in patients with pulmonary embolism, Outcome 4 Incidence of major haemorrhagic episodes (during initial treatment).

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 3 LMWH versus UFH in patients with pulmonary embolism

Outcome: 4 Incidence of major haemorrhagic episodes (during initial treatment)

tudy or subgroup	LMWH n/N	UFH n/N	Peto Odds Ratio Peto,Fixed,95% Cl	Weight	Peto Odds Ratio Peto,Fixed,95% Cl
droparin versus unfractionate	ed heparin				
hery 1992	0/35	2/33		66.9 %	0.12[0.01, 2.02]
ototal (95% CI)	35	33		66.9 %	0.12 [0.01, 2.02]
events: 0 (LMWH), 2 (UFH)					
erogeneity: not applicable					
for overall effect: Z = 1.47 (P	= 0.14)				
oxaparin versus unfractionate	d heparin				
rez de Llano 2003	1/29	0/21		33.1 %	5.61 [0.11, 297.44]
ototal (95% CI)	29	21		33.1 %	5.61 [0.11, 297.44]
events: I (LMWH), 0 (UFH)					
erogeneity: not applicable					
for overall effect: Z = 0.85 (P	= 0.39)				
Iteparin versus unfractionated	l heparin				
leyer 1995	0/29	0/31			Not estimable
ototal (95% CI)	29	31			Not estimable
events: 0 (LMWH), 0 (UFH)					
erogeneity: not applicable					
for overall effect: not applicab	le				
al (95% CI)	93	85	-	100.0 %	0.44 [0.04, 4.29]
events: I (LMWH), 2 (UFH)					
erogeneity: Chi ² = 2.37, df =	$ (P = 0.12); ^2 =$	=58%			
for overall effect: Z = 0.71 (P	= 0.48)				
for subgroup differences: Chi ²	2 = 2.37, df = 1	$(P = 0.12), I^2 = 58\%$			

0.001 0.01 0.1 1 10 100 1000 Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous III thromboembolism (Review)

Analysis 3.5. Comparison 3 LMWH versus UFH in patients with pulmonary embolism, Outcome 5 Overall mortality at end of follow-up.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 3 LMWH versus UFH in patients with pulmonary embolism

Outcome: 5 Overall mortality at end of follow-up

Study or subgroup	LMWH	UFH	Peto Odds Ratio	Weight	Peto Odds Ratio
	n/N	n/N	Peto,Fixed,95% Cl		Peto,Fixed,95% Cl
I Nadroparin versus unfractiona	ated heparin				
Thery 1992	1/35	1/33		66.9 %	0.94 [0.06, 15.40]
Subtotal (95% CI)	35	33	-	66.9 %	0.94 [0.06, 15.40]
Total events: (LMWH), (UFF	H)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 0.04$	(P = 0.97)				
2 Enoxaparin versus unfractiona	ited heparin				
P rez de Llano 2003	1/29	0/21		33.1 %	5.61 [0.11, 297.44]
Subtotal (95% CI)	29	21		33.1 %	5.61 [0.11, 297.44]
Total events: (LMWH), 0 (UFF	H)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 0.85$	(P = 0.39)				
3 Dalteparin versus unfractionat	ted heparin				
Meyer 1995	0/29	0/31			Not estimable
Subtotal (95% CI)	29	31			Not estimable
Total events: 0 (LMWH), 0 (UFF	H)				
Heterogeneity: not applicable					
Test for overall effect: not applic	able				
Total (95% CI)	93	85	-	100.0 %	1.70 [0.17, 16.71]
Total events: 2 (LMWH), 1 (UFF	H)				
Heterogeneity: Chi ² = 0.52, df =	= I (P = 0.47); I ² =	=0.0%			
Test for overall effect: $Z = 0.46$	(P = 0.65)				
Test for subgroup differences: C	$hi^2 = 0.52, df = 1$	$(P = 0.47), I^2 = 0.0\%$			

0.001 0.01 0.1 1 10 100 1000 Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 112 thromboembolism (Review)

Analysis 4.1. Comparison 4 LMWH versus UFH in patients with venous thromboembolism and malignant disease, Outcome 1 Mortality at the end of follow-up.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 4 LMWH versus UFH in patients with venous thromboembolism and malignant disease

Outcome: I Mortality at the end of follow-up

Study or subgroup	LMWH n/N	UFH n/N	Peto Odds Ratio Peto,Fixed,95% Cl	Weight	Peto Odds Ratio Peto,Fixed,95% Cl
I Mortality in patients with ma	lignant disease				
Columbus 1997	20/119	27/113	-	55.1 %	0.65 [0.34, 1.22]
Hull 1992	6/47	13/49		22.6 %	0.42 [0.16, 1.15]
Lindmarker 1994	2/7	2/9	· · · · · ·	4.6 %	1.37 [0.15, 12.50]
Lopaciuk 1992	0/7	0/2			Not estimable
Prandoni 1992	1/15	8/18		9.8 %	0.16 [0.03, 0.72]
Simonneau 1997	2/26	4/34		7.9 %	0.64 [0.12, 3.47]
Subtotal (95% CI)	221	225	•	100.0 %	0.53 [0.33, 0.85]
Total events: 31 (LMWH), 54 ((UFH)				
Heterogeneity: $Chi^2 = 3.78$, df	$r = 4 (P = 0.44); l^2 =$	=0.0%			
Test for overall effect: $Z = 2.63$	B (P = 0.0086)				
2 Mortality in patients with ma	lignant disease in tri	al with adequate co	oncealment		
Columbus 1997	20/119	27/113	-	57.8 %	0.65 [0.34, 1.22]
Hull 1992	6/47	13/49		23.6 %	0.42 [0.16, 1.15]
Lopaciuk 1992	0/7	0/2			Not estimable
Prandoni 1992	1/15	8/18		10.3 %	0.16 [0.03, 0.72]
Simonneau 1997	2/26	4/34		8.3 %	0.64 [0.12, 3.47]
Subtotal (95% CI)	214	216	•	100.0 %	0.51 [0.31, 0.82]
Total events: 29 (LMWH), 52 ((UFH)				
Heterogeneity: $Chi^2 = 3.03$, df	$= 3 (P = 0.39); I^2 =$	-1%			
Test for overall effect: $Z = 2.75$	5 (P = 0.0059)				
			0.01 0.1 1 10 100		
			Favours LMWH Favours UFH		

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 113

thromboembolism (Review)

Analysis 5.1. Comparison 5 LMWH versus UFH in patients with venous thromboembolism without malignant disease, Outcome I Mortality at the end of follow-up.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 5 LMWH versus UFH in patients with venous thromboembolism without malignant disease

Outcome: I Mortality at the end of follow-up

Study or subgroup	LMWH n/N	UFH n/N	Peto Odds Ratio Peto,Fixed,95% Cl	Weight	Peto Odds Ratio Peto,Fixed,95% Cl
I Mortality in patients without	malignant disease				
Columbus 1997	16/391	12/398	+	39.5 %	1.37 [0.64, 2.91]
Hull 1992	4/166	8/170		17.0 %	0.51 [0.16, 1.63]
Lindmarker 1994	0/94	1/94	·	1.5 %	0.14 [0.00, 6.82]
Lopaciuk 1992	0/67	1/70	·	1.5 %	0.14 [0.00, 7.13]
Prandoni 1992	5/70	4/67	-	12.4 %	1.21 [0.31, 4.65]
Simonneau 1997	10/278	10/274	+	28.2 %	0.99 [0.40, 2.40]
Subtotal (95% CI) Total events: 35 (LMWH), 36 (I Heterogeneity: $Chi^2 = 3.97$, df Test for overall effect: $Z = 0.11$	= 5 (P = 0.55); I ² =	1073	•	100.0 %	0.97 [0.61, 1.56]
2 Mortality in patients without		trials with adequate o	concealment		
Columbus 1997	6/39	12/398	+	40.1 %	1.37 [0.64, 2.91]
Hull 1992	4/166	8/170		17.2 %	0.51 [0.16, 1.63]
Lopaciuk 1992	0/67	1/70	·	1.5 %	0.14 [0.00, 7.13]
Prandoni 1992	5/70	4/67	-	12.6 %	1.21 [0.31, 4.65]
Simonneau 1997	10/278	10/274	+	28.6 %	0.99 [0.40, 2.40]
Subtotal (95% CI)	972	979	+	100.0 %	1.00 [0.62, 1.62]
Total events: 35 (LMWH), 35 (Heterogeneity: Chi ² = 2.98, df Test for overall effect: $Z = 0.01$	= 4 (P = 0.56); l ² =	=0.0%			

Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 114 thromboembolism (Review)

Analysis 6.1. Comparison 6 LMWH versus subcutaneous UFH in patients with venous thromboembolism, Outcome I Incidence of recurrent venous thromboembolism at the end of follow-up.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 6 LMWH versus subcutaneous UFH in patients with venous thromboembolism

Outcome: I Incidence of recurrent venous thromboembolism at the end of follow-up

Study or subgroup	LMWH	Subcutaneous UFH	Peto Odds Ratio	Weight	Peto Odds Ratio
	n/N	n/N	Peto,Fixed,95% Cl		Peto,Fixed,95% CI
Leizorovicz 2011	7/269	3/268		24.7 %	2.25 [0.64, 7.85]
Lopaciuk 1992	0/74	3/72		7.4 %	0.13[0.01, 1.25]
Prandoni 2004	14/360	14/360	+	67.8 %	1.00 [0.47, 2.13]
Total (95% CI)	703	700	+	100.0 %	1.05 [0.56, 1.95]
Total events: 21 (LMWH), 20 (Subcutaneou	is UFH)			
Heterogeneity: $Chi^2 = 4$.	72, df = 2 (P = 0.0	9); l ² =58%			
Test for overall effect: Z =	= 0.15 (P = 0.88)				
Test for subgroup differer	nces: Not applicabl	e			
			0.01 0.1 1 10 10	0	
			Favours LMWH Favours UFH		

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 115 thromboembolism (Review)

Analysis 6.2. Comparison 6 LMWH versus subcutaneous UFH in patients with venous thromboembolism, Outcome 2 Incidence of major haemorrhagic episodes (during initial treatment).

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 6 LMWH versus subcutaneous UFH in patients with venous thromboembolism

Outcome: 2 Incidence of major haemorrhagic episodes (during initial treatment)

Study or subgroup	LMWH	Subcutaneous UFH	Peto Odds Ratio	Weight	Peto Odds Ratio
	n/N	n/N	Peto,Fixed,95% Cl	-	Peto,Fixed,95% Cl
Faivre 1988	0/33	3/35		6.9 %	0.13[0.01, 1.34]
Leizorovicz 2011	18/269	15/268	+	74.1 %	1.21 [0.60, 2.44]
Lopaciuk 1992	0/74	1/72	·	2.4 %	0.13 [0.00, 6.64]
Prandoni 2004	3/360	4/360		16.6 %	0.75 [0.17, 3.32]
Total (95% CI) Total events: 21 (LMWH) Heterogeneity: $Chi^2 = 4$.	,	,	•	100.0 %	0.91 [0.50, 1.67]
Test for overall effect: Z = Test for subgroup differen	· · · · ·	le			

0.0010.010.11 I 10 100 1000 Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 116 thromboembolism (Review)

Analysis 6.3. Comparison 6 LMWH versus subcutaneous UFH in patients with venous thromboembolism, Outcome 3 Overall mortality at the end of follow-up.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 6 LMWH versus subcutaneous UFH in patients with venous thromboembolism

Outcome: 3 Overall mortality at the end of follow-up

Study or subgroup	LMWH	Subcutaneous UFH	Peto Odds Ratio	Weight	Peto Odds Ratio
	n/N	n/N	Peto,Fixed,95% Cl		Peto,Fixed,95% Cl
Leizorovicz 2011	31/269	17/268	-	64.4 %	1.89 [1.04, 3.41]
Lopaciuk 1992	0/74	1/72	·	1.5 %	0.13 [0.00, 6.64]
Prandoni 2004	12/360	12/360	+	34.2 %	1.00 [0.44, 2.26]
Total (95% CI)	703	700	•	100.0 %	1.46 [0.91, 2.35]
Total events: 43 (LMWH)), 30 (Subcutaneou	s UFH)			
Heterogeneity: $Chi^2 = 3.0$	00, df = 2 (P = 0.2	2); ² =33%			
Test for overall effect: Z =	= 1.56 (P = 0.12)				
Test for subgroup differer	nces: Not applicable	e			

0.001 0.01 0.1 1 10 100 1000 Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 117 thromboembolism (Review)

Analysis 7.1. Comparison 7 LMWH versus intravenous UFH in patients with venous thromboembolism, Outcome I Incidence of recurrent venous thromboembolism at the end of follow-up.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 7 LMWH versus intravenous UFH in patients with venous thromboembolism

Outcome: I Incidence of recurrent venous thromboembolism at the end of follow-up

Pet Odds Rati Peto,Fixed,95% (Weight	Peto Odds Ratio Peto,Fixed,95% Cl	dy or subgroup LMWH Intravenous UFH Odds Ratio n/N n/N Peto,Fixed,95% Cl		Study or subgroup
0.92 [0.34, 2.46	4.7 %	-	13/196	6/98	Belcaro 1999
0.31 [0.15, 0.63	8.8 %	-#-	24/375	7/388	Breddin 2001
1.09 [0.62, 1.90	14.5 %	+	25/511	27/510	Columbus 1997
0.87 [0.37, 2.05	6.1 %		12/205	10/195	Decousus 1998
0.36 [0.05, 2.70	1.1 %		3/30	1/29	Findik 2002
0.13 [0.00, 6.49	0.3 %	·	1/39	0/41	Goldhaber 1998
0.42 [0.18, 1.01	5.9 %		15/273	6/265	Harenberg 2000a
0.42 [0.17, 1.01	5.9 %		15/219	6/213	Hull 1992
Not estimab			0/110	0/	Kakkar 2003
0.50 [0.10, 2.53	1.7 %	<u> </u>	4/124	2/125	Kirchmaier 1998
0.79 [0.38, 1.65	8.4 %	-	17/198	14/202	Koopman 1996
0.77 [0.37, 1.62	8.3 %	-	17/253	13/247	Levine 1996
1.71 [0.42, 7.02	2.3 %		3/103	5/101	Lindmarker 1994
0.90 [0.42, 1.92	7.9 %	-	11/290	21/610	Merli 2001
Not estimab			0/3	0/29	Meyer 1995
0.48 [0.18, 1.26	4.8 %	<u> </u>	12/85	6/85	Prandoni 1992
1.09 [0.17, 6.98	1.3 %		2/21	3/29	P rez de Llano 2003
0.76 [0.43, 1.35	13.8 %	+	27/593	22/627	Riess 2003
0.13 [0.01, 1.28	0.9 %		3/67	0/67	Simonneau 1993
0.84 [0.26, 2.77	3.2 %		6/308	5/304	Simonneau 1997
Not estimab			0/33	0/35	Thery 1992
0.69 [0.56, 0.86	100.0 %	•	,	df = 17 (P = 0.45) 8 (P = 0.00072)	otal (95% CI) tal events: 154 (LMWH), 2 eterogeneity: Chi ² = 17.06, st for overall effect: Z = 3: st for subgroup differences

0.0010.010.1101001000 Favours LMWH Favours iv UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 118 thromboembolism (Review)

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Analysis 7.2. Comparison 7 LMWH versus intravenous UFH in patients with venous thromboembolism, Outcome 2 Incidence of major haemorrhagic episodes (during initial treatment).

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 7 LMWH versus intravenous UFH in patients with venous thromboembolism

Outcome: 2 Incidence of major haemorrhagic episodes (during initial treatment)

Pet Odds Rat Peto,Fixed,95% (Weight	Peto Odds Ratio Peto,Fixed,95% Cl	Intravenous UFH n/N	LMWH n/N	Study or subgroup
Not estimab			0/196	0/98	Belcaro 1999
1.26 [0.49, 3.19	15.7 %	-	8/511	10/510	Columbus 1997
0.92 [0.33, 2.57	12.8 %	-	8/205	7/195	Decousus 1998
0.15 [0.01, 2.39	1.8 %		2/133	0/120	Fiessinger 1996
Not estimab			0/30	0/29	Findik 2002
0.40 [0.14, 1.10	13.0 %		11/273	4/265	Harenberg 2000a
0.19 [0.06, 0.59	10.4 %		11/219	1/213	Hull 1992
7.39 [0.46, 118.89	1.8 %		0/110	2/111	Kakkar 2003
0.30 [0.05, 1.77	4.4 %	- _	4/131	1/128	Kirchmaier 1998
0.50 [0.05, 4.85	2.6 %		2/198	1/202	Koopman 1996
1.70 [0.42, 6.87	7.0 %		3/253	5/247	Levine 1996
Not estimab			0/103	0/101	Lindmarker 1994
0.15 [0.00, 7.64	0.9 %	·	1/131	0/117	Luomanmaki 1996
Not estimab			0/31	0/29	Meyer 1995
0.48 [0.09, 2.43	5.2 %		4/81	2/85	Ninet 1991
0.36 [0.05, 2.61	3.5 %		3/85	1/85	Prandoni 1992
5.61 [0.11, 297.44	0.9 %		0/21	1/29	P rez de Llano 2003
0.81 [0.27, 2.41	11.4 %		7/593	6/627	Riess 2003
Not estimab			0/67	0/67	Simonneau 1993
0.61 [0.15, 2.46	7.0 %		5/308	3/304	Simonneau 1997
0.12 [0.01, 2.02	1.7 %		2/33	0/35	Thery 1992
0.62 [0.43, 0.90	100.0 %	•	3712	3597	otal (95% CI)
				df = 15 (P = 0.26) 61 (P = 0.012)	tal events: 44 (LMWH), 71 eterogeneity: $Chi^2 = 18.06$, st for overall effect: $Z = 2.5$ st for subgroup differences

Favours LMWH Favours iv UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 119 thromboembolism (Review)

Analysis 7.3. Comparison 7 LMWH versus intravenous UFH in patients with venous thromboembolism, Outcome 3 Overall mortality at the end of follow-up.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 7 LMWH versus intravenous UFH in patients with venous thromboembolism

Outcome: 3 Overall mortality at the end of follow-up

Pe Odds Rat Peto,Fixed,95%	Weight	Peto Odds Ratio Peto,Fixed,95% Cl	udy or subgroup LMWH Intravenous UFH n/N n/N		ogroup LMWH Intravenous UFH Odds Ratio		Study or subgroup
0.79 [0.32, .9]	5.0 %		1/375	9/388	Breddin 2001		
0.92 [0.57, 1.47	18.0 %	+	39/511	36/510	Columbus 1997		
0.97 [0.60, 1.58	17.0 %	+	43/205	40/195	Decousus 1998		
Not estimat			0/30	0/29	Findik 2002		
Not estimat			0/39	0/41	Goldhaber 1998		
0.42 [0.18, 1.01	5.2 %		15/273	6/265	Harenberg 2000a		
0.48 [0.23, 1.00	7.5 %		21/219	10/213	Hull 1992		
0.95 [0.13, 6.90	1.0 %		2/85	2/89	Kakkar 2003		
0.78 [0.30, 2.03	4.3 %		10/124	8/125	Kirchmaier 1998		
0.85 [0.40, 1.78	7.2 %	+	16/198	14/202	Koopman 1996		
0.65 [0.30, 1.40	6.8 %		17/253	11/247	Levine 1996		
0.68 [0.12, 3.99	1.3 %	<u> </u>	3/103	2/101	Lindmarker 1994		
0.31 [0.05, 1.82	1.3 %		4/116	1/110	Luomanmaki 1996		
0.95 [0.42, 2.15	5.9 %	-	9/290	18/610	Merli 2001		
Not estimat			0/31	0/29	Meyer 1995		
0.48 [0.18, 1.26	4.2 %		12/85	6/85	Prandoni 1992		
5.61 [0.11, 297.44	0.3 %		0/21	1/29	P rez de Llano 2003		
0.65 [0.30, 1.39	6.8 %		16/593	11/627	Riess 2003		
1.51 [0.25, 8.96	1.3 %	<u>_</u>	2/67	3/67	Simonneau 1993		
0.86 [0.39, 1.89	6.4 %	-	14/308	12/304	Simonneau 1997		
0.94 [0.06, 15.40	0.5 %		1/33	1/35	Thery 1992		
0.77 [0.63, 0.93	100.0 %	•	· · · · · · · · · · · · · · · · · · ·	Hf = 17 (P = 0.93); 3 (P = 0.0085)	btal (95% CI) tal events: 191 (LMWH), 2 eterogeneity: Chi ² = 9.17, o st for overall effect: Z = 2.0 st for subgroup differences		

0.001 0.01 0.1 1 10 100 1000 Favours LMWH Favours iv UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 120 thromboembolism (Review)

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Analysis 8.1. Comparison 8 LMWH versus UFH: all randomised controlled trials with adequate concealment of allocation, Outcome 1 Incidence of recurrent venous thromboembolism during initial treatment.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 8 LMWH versus UFH: all randomised controlled trials with adequate concealment of allocation

Outcome: I Incidence of recurrent venous thromboembolism during initial treatment

Study or subgroup	LMWH n/N	UFH n/N	Peto Odds Ratio Peto,Fixed,95% Cl	Weight	Peto Odds Ratio Peto,Fixed,95% Cl
I Nadroparin versus unfraction		11/1 N	1 eto,1 ixed,75% Ci		1 eto,1 1xed,75% CI
Koopman 1996	4/202	5/198		7.8 %	0.78 [0.21, 2.92]
Lopaciuk 1992	0/74	1/72	·	0.9 %	0.13 [0.00, 6.64]
·					
Prandoni 1992	1/85	4/85		4.3 %	0.29 [0.05, 1.72]
Subtotal (95% CI)	361	355	•	13.0 %	0.50 [0.18, 1.39]
Total events: 5 (LMWH), 10 (L Heterogeneity: $Chi^2 = 1.23$, df	·	0.0%			
Test for overall effect: $Z = 1.33$	· · · ·	-0.076			
2 Tinzaparin versus unfractiona	. ,				
Simonneau 1997	3/304	2/308	_ 	4.4 %	1.52 [0.26, 8.80]
Subtotal (95% CI)	304	308	-	4.4 %	1.52 [0.26, 8.80]
Total events: 3 (LMWH), 2 (UF	FH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 0.46$	· /				
3 Enoxaparin versus unfraction		0,005		0.5.0/	
Decousus 1998	3/195	8/205	•	9.5 %	0.41 [0.13, 1.37]
Levine 1996	7/247	12/253		16.2 %	0.59 [0.24, 1.48]
Simonneau 1993	0/67	2/67		1.8 %	0.13 [0.01, 2.15]
Subtotal (95% CI)	509	525	•	27.4 %	0.48 [0.24, 0.96]
Total events: 10 (LMWH), 22 (· · ·				
Heterogeneity: $Chi^2 = 1.08$, df		0.0%			
Test for overall effect: $Z = 2.06$	` '				
4 Reviparin versus unfractionat					
Columbus 1997	16/510	15/511	T	26.6 %	1.07 [0.52, 2.19]
Subtotal (95% CI)	510	511	+	26.6 %	1.07 [0.52, 2.19]
Total events: 16 (LMWH), 15 ((UFH)				
Heterogeneity: not applicable	(D - 0.05)				
Test for overall effect: Z = 0.19 5 Certoparin versus unfraction	· /				
5 Certoparin versus uniraction	lated heparin				
			0.001 0.01 0.1 1 10 100 1000		
			Favours LMWH Favours UFH		

(Continued . . .)

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 121 thromboembolism (Review)

Study or subgroup	LMWH n/N	UFH n/N	Peto Odds Ratio Peto,Fixed,95% Cl	Weight	(Continued) Peto Odds Ratio Peto,Fixed,95% Cl
Kirchmaier 1998	1/128	3/131		3.5 %	0.37 [0.05, 2.67]
Riess 2003	14/627	15/593	+	25.0 %	0.88 [0.42, 1.84]
Subtotal (95% CI)	755	724	•	28.5 %	0.79 [0.40, 1.58]
Total events: 15 (LMWH), 18 ((UFH)				
Heterogeneity: Chi ² = 0.64, df	= (P = 0.42); ² =	=0.0%			
Test for overall effect: $Z = 0.66$	6 (P = 0.51)				
Total (95% CI)	2439	2423	•	100.0 %	0.72 [0.50, 1.05]
Total events: 49 (LMWH), 67 ((UFH)				
Heterogeneity: $Chi^2 = 6.72$, df	= 9 (P = 0.67); l ² =	=0.0%			
Test for overall effect: $Z = 1.72$	e (P = 0.085)				
Test for subgroup differences: (Chi ² = 3.76, df = 4	$(P = 0.44), I^2 = 0.0\%$			

0.001 0.01 0.1 1 10 100 1000

Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 122 thromboembolism (Review)

Analysis 8.2. Comparison 8 LMWH versus UFH: all randomised controlled trials with adequate concealment of allocation, Outcome 2 Incidence of recurrent venous thromboembolism at the end of followup.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 8 LMWH versus UFH: all randomised controlled trials with adequate concealment of allocation

Outcome: 2 Incidence of recurrent venous thromboembolism at the end of follow-up

Study or subgroup	LMWH	UFH	Peto Odds Ratio	Weight	Peto Odds Ratio	
,	n/N	n/N	Peto,Fixed,95% Cl	0	Peto,Fixed,95% Cl	
I Nadroparin versus unfraction	nated heparin					
Koopman 1996	14/202	17/198	-	9.9 %	0.79 [0.38, 1.65]	
Lopaciuk 1992	0/74	3/72		1.0 %	0.13 [0.01, 1.25]	
Prandoni 1992	6/85	12/85		5.6 %	0.48 [0.18, 1.26]	
Prandoni 2004	14/360	15/360	+	9.6 %	0.93 [0.44, 1.96]	
Subtotal (95% CI)	721	715	•	26.2 %	0.70 [0.45, 1.10]	
Total events: 34 (LMWH), 47 (Heterogeneity: Chi ² = 3.41, df Test for overall effect: Z = 1.53 2 Tinzaparin versus unfractiona Hull 1992	$f = 3 (P = 0.33); I^2 = 3 (P = 0.12)$	=12% 15/219	-	6.9 %	0.42 [0.17, 1.01]	
	5/204	6/308	_	3.7 %	0.84 [0.26, 2.77]	
Simonneau 1997	5/304			5.7 70	0.01[0.20, 2.77]	
Simonneau 1997 Subtotal (95% CI) Total events: 11 (LMWH), 21 (. ,	527	•	10.7 %	0.54 [0.26, 1.08]	
Subtotal (95% CI)	517 (UFH) F = I (P = 0.35); I ² = 4 (P = 0.083)		•	10.7 % 0.3 %	0.54 [0.26, 1.08] 0.13 [0.00, 6.65]	
Subtotal (95% CI) Total events: 11 (LMWH), 21 (Heterogeneity: Chi ² = 0.86, df Test for overall effect: Z = 1.74 3 Ardeparin versus unfractional	517 (UFH) $f = 1 (P = 0.35); I^2 = 4 (P = 0.083)$ ated heparin 0/41 41 FH) 1 (P = 0.31)	=0.0%	•			
 Subtotal (95% CI) Total events: 11 (LMWH), 21 (1) Heterogeneity: Chi² = 0.86, df Test for overall effect: Z = 1.74 3 Ardeparin versus unfractional Goldhaber 1998 Subtotal (95% CI) Total events: 0 (LMWH), 1 (UI) Heterogeneity: not applicable Test for overall effect: Z = 1.01 	517 (UFH) $f = 1 (P = 0.35); I^2 = 4 (P = 0.083)$ ated heparin 0/41 41 FH) 1 (P = 0.31)	=0.0%		0.3 %	0.13 [0.00, 6.65]	
 Subtotal (95% CI) Total events: 11 (LMWH), 21 (1) Heterogeneity: Chi² = 0.86, df Test for overall effect: Z = 1.74 3 Ardeparin versus unfractiona Goldhaber 1998 Subtotal (95% CI) Total events: 0 (LMWH), 1 (UI) Heterogeneity: not applicable Test for overall effect: Z = 1.01 4 Enoxaparin versus unfractional 	517 (UFH) $F = 1 (P = 0.35); I^2 = 4 (P = 0.083)$ ated heparin 0/4 I 41 FH) I (P = 0.3 I) nated heparin	=0.0% 1/40 40		0.3 % 0.3 %	0.13 [0.00, 6.65] 0.13 [0.00, 6.65]	
 Subtotal (95% CI) Total events: 11 (LMWH), 21 (1) Heterogeneity: Chi² = 0.86, df Test for overall effect: Z = 1.74 3 Ardeparin versus unfractional Goldhaber 1998 Subtotal (95% CI) Total events: 0 (LMWH), 1 (UI) Heterogeneity: not applicable Test for overall effect: Z = 1.01 4 Enoxaparin versus unfraction Decousus 1998 	517 (UFH) $f = 1 (P = 0.35); I^2 = 4 (P = 0.083)$ ated heparin 0/41 41 FH) 1 (P = 0.31) mated heparin 10/195	=0.0% 1/40 40 12/205		0.3 % 0.3 % 7.2 %	0.13 [0.00, 6.65] 0.13 [0.00, 6.65] 0.87 [0.37, 2.05]	
 Subtotal (95% CI) Total events: 11 (LMWH), 21 Heterogeneity: Chi² = 0.86, df Test for overall effect: Z = 1.74 3 Ardeparin versus unfractional Goldhaber 1998 Subtotal (95% CI) Total events: 0 (LMWH), 1 (UI Heterogeneity: not applicable Test for overall effect: Z = 1.01 4 Enoxaparin versus unfraction Decousus 1998 Levine 1996 	517 (UFH) $F = 1 (P = 0.35); I^2 = 4 (P = 0.083)$ ated heparin 0/41 41 FH) 1 (P = 0.31) nated heparin 10/195 13/247	=0.0% 1/40 40 12/205 17/253		0.3 % 0.3 % 7.2 % 9.8 %	0.13 [0.00, 6.65] 0.13 [0.00, 6.65] 0.87 [0.37, 2.05] 0.77 [0.37, 1.62]	

Favours LMWH Favours UFH

(Continued ...)

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 123 thromboembolism (Review)

Study or subgroup	LMWH n/N	UFH n/N	Peto Odds Ratio Peto,Fixed,95% Cl	Weight	(Continued) Peto Odds Ratio Peto,Fixed,95% Cl
Heterogeneity: $Chi^2 = 2.55$, df	= 3 (P = 0.47); l ² =	=0.0%			
Test for overall effect: $Z = 1.07$	(P = 0.29)				
5 Reviparin versus unfractionat	ed heparin				
Columbus 1997	27/510	25/510	+	17.1 %	1.08 [0.62, 1.89]
Subtotal (95% CI)	510	510	•	17.1 %	1.08 [0.62, 1.89]
Total events: 27 (LMWH), 25 (UFH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 0.28$	(P = 0.78)				
6 Certoparin versus unfraction	ated heparin				
Kirchmaier 1998	2/125	4/124		2.0 %	0.50 [0.10, 2.53]
Riess 2003	22/627	27/593	+	16.3 %	0.76 [0.43, 1.35]
Subtotal (95% CI)	752	717	•	18.3 %	0.73 [0.42, 1.25]
Total events: 24 (LMWH), 31 (UFH)				
Heterogeneity: $Chi^2 = 0.23$, df	$= (P = 0.63); ^2 =$	=0.0%			
Test for overall effect: $Z = 1.15$	(P = 0.25)				
Total (95% CI)	3660	3324	•	100.0 %	0.76 [0.60, 0.96]
Total events: 140 (LMWH), 16	8 (UFH)				
Heterogeneity: $Chi^2 = 10.49$, c	lf = 13 (P = 0.65); I	2 =0.0%			
Test for overall effect: $Z = 2.35$	(P = 0.019)				
Test for subgroup differences: ($Chi^2 = 3.44, df = 5$	(P = 0.63), I ² =0.0%			

0.001 0.01 0.1 1 10 100 1000

Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 124 thromboembolism (Review)

Analysis 8.3. Comparison 8 LMWH versus UFH: all randomised controlled trials with adequate concealment of allocation, Outcome 3 Incidence of recurrent venous thromboembolism at 3 months' followup.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 8 LMWH versus UFH: all randomised controlled trials with adequate concealment of allocation

Outcome: 3 Incidence of recurrent venous thromboembolism at 3 months' follow-up

Study or subgroup	LMWH	UFH	Peto Odds Ratio	Weight	Peto Odds Ratio
,	n/N	n/N	Peto,Fixed,95% Cl	0	Peto,Fixed,95% Cl
I Nadroparin versus unfractior	nated heparin				
Koopman 1996	8/202	10/198		7.9 %	0.78 [0.30, 2.00]
Lopaciuk 1992	0/74	3/72		1.4 %	0.13 [0.01, 1.25]
Prandoni 1992	4/85	7/85		4.8 %	0.56 [0.17, 1.89]
Prandoni 2004	14/360	15/360	-	12.8 %	0.93 [0.44, 1.96]
Subtotal (95% CI)	721	715	•	26.8 %	0.73 [0.44, 1.22]
Total events: 26 (LMWH), 35 (UFH)				
Heterogeneity: $Chi^2 = 2.85$, df	= 3 (P = 0.42); l ² =	=0.0%			
Test for overall effect: $Z = 1.21$	· /				
2 Tinzaparin versus unfractiona					
Hull 1992	6/213	15/219		9.2 %	0.42 [0.17, 1.01]
Simonneau 1997	5/304	6/308		5.0 %	0.84 [0.26, 2.77]
Subtotal (95% CI)	517	527	•	14.2 %	0.54 [0.26, 1.08]
Total events: 11 (LMWH), 21 (UFH)				
Heterogeneity: $Chi^2 = 0.86$, df	$= 1 (P = 0.35); I^2 =$	=0.0%			
Test for overall effect: $Z = 1.74$,				
3 Enoxaparin versus unfraction					
Decousus 1998	10/195	12/205		9.6 %	0.87 [0.37, 2.05]
Levine 1996	13/247	17/253		13.0 %	0.77 [0.37, 1.62]
Merli 2001	21/610	11/290		12.4 %	0.90 [0.42, 1.92]
Simonneau 1993	0/67	3/67		1.4 %	0.13 [0.01, 1.28]
Subtotal (95% CI)	1119	815	+	36.3 %	0.79 [0.51, 1.22]
Total events: 44 (LMWH), 43 (UFH)				
Heterogeneity: $Chi^2 = 2.55$, df	= 3 (P = 0.47); I ² =	=0.0%			
Test for overall effect: $Z = 1.07$	(P = 0.29)				
4 Reviparin versus unfractionat					
Columbus 1997	27/510	25/511		22.7 %	1.09 [0.62, 1.90]
Subtotal (95% CI) Total events: 27 (LMWH), 25 (510 (UFH)	511	+	22.7 %	1.09 [0.62, 1.90]
			0.005 0.1 1 10 200		
			Favours LMWH Favours UFH		

(Continued . . .)

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 125 thromboembolism (Review)

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Study or subgroup	LMWH n/N	UFH n/N			lds R	Peto atio ,95% Cl		Weight	(Continued) Peto Odds Ratio Peto,Fixed,95% Cl
Heterogeneity: not applicable									
Test for overall effect: $Z = 0.2$	9 (P = 0.77)								
Total (95% CI)	2867	2568			•			100.0 %	0.79 [0.60, 1.02]
Total events: 108 (LMWH), 1	24 (UFH)								
Heterogeneity: $Chi^2 = 8.77$, c	$f = 10 (P = 0.55); I^2$	=0.0%							
Test for overall effect: $Z = 1.7$	78 (P = 0.075)								
Test for subgroup differences:	$Chi^2 = 2.5I, df = 3$ ($P = 0.47$), $I^2 = 0.0\%$							
			0.005	0.1	I	10	200		
			Favours	s LMWH		Favours	UFH		

Analysis 8.4. Comparison 8 LMWH versus UFH: all randomised controlled trials with adequate concealment of allocation, Outcome 4 Reduction in thrombus size (pre- and post-treatment venograms).

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 8 LMWH versus UFH: all randomised controlled trials with adequate concealment of allocation

Outcome: 4 Reduction in thrombus size (pre- and post-treatment venograms)

Study or subgroup	LMWH	UFH	Peto Odds Ratio(Non-event)	Weight	Peto Odds Ratio(Non-event)	
	n/N	n/N	Peto,Fixed,95% CI	-	Peto,Fixed,95% Cl	
I Nadroparin versus unfraction	nated heparin					
Lopaciuk 1992	45/68	32/66		18.2 %	0.49 [0.25, 0.96]	
Prandoni 1992	50/83	36/85		23.3 %	0.49 [0.27, 0.90]	
Subtotal (95% CI)	151	151	•	41.4 %	0.49 [0.31, 0.77]	
Total events: 95 (LMWH), 68 (UFH)					
Heterogeneity: $Chi^2 = 0.00$, df	$= (P = 0.99); ^2 =$	0.0%				
Test for overall effect: $Z = 3.10$	(P = 0.0019)					
2 Ardeparin versus unfractiona	ted heparin					
Goldhaber 1998	31/39	21/36		8.9 %	0.37 [0.14, 0.99]	
Subtotal (95% CI)	39	36	-	8.9 %	0.37 [0.14, 0.99]	
Total events: 31 (LMWH), 21 (UFH)					
Heterogeneity: not applicable						
			0.01 0.1 1 10 100			
			Favours LMWH Favours UFH			

(Continued . . .)

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 126 thromboembolism (Review)

					(Continued)
Study or subgroup	LMWH	UFH	Peto Odds Ratio(Non-event)	Weight	Peto Odds Ratio(Non-event)
	n/N	n/N	Peto,Fixed,95% CI	-	Peto,Fixed,95% Cl
Test for overall effect: Z = 1.97	7 (P = 0.049)				
3 Enoxparin versus unfractiona	ted heparin				
Simonneau 1993	35/60	18/57		16.1 %	0.34 [0.17, 0.71]
Subtotal (95% CI)	60	57	•	16.1 %	0.34 [0.17, 0.71]
Total events: 35 (LMWH), 18 ((UFH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 2.89$	P (P = 0.0038)				
4 Certoparin versus unfraction	ated heparin				
Kirchmaier 1998	55/128	42/131	-	33.6 %	0.63 [0.38, 1.04]
Subtotal (95% CI)	128	131	•	33.6 %	0.63 [0.38, 1.04]
Total events: 55 (LMWH), 42 ((UFH)				
Heterogeneity: not applicable					
Test for overall effect: Z = 1.81	(P = 0.070)				
Total (95% CI)	378	375	•	100.0 %	0.49 [0.37, 0.66]
Total events: 216 (LMWH), 14	9 (UFH)				
Heterogeneity: Chi ² = 2.17, df	= 4 (P = 0.70); l ² =	=0.0%			
Test for overall effect: Z = 4.79	P (P < 0.00001)				
Test for subgroup differences: (Chi ² = 2.17, df = 3	(P = 0.54), I ² =0.0%			

0.01 0.1 1 10 100

Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 127 thromboembolism (Review)

Analysis 8.5. Comparison 8 LMWH versus UFH: all randomised controlled trials with adequate concealment of allocation, Outcome 5 Incidence of major haemorrhagic episodes (during initial treatment).

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 8 LMWH versus UFH: all randomised controlled trials with adequate concealment of allocation

Outcome: 5 Incidence of major haemorrhagic episodes (during initial treatment)

Pet Odds Rat	Weight	Peto Odds Ratio	UFH	LMWH	Study or subgroup
Peto,Fixed,95% (Peto,Fixed,95% CI	n/N	n/N	
				ated heparin	I Nadroparin versus unfraction
0.50 [0.05, 4.85	3.2 %		2/198	1/202	Koopman 1996
0.13 [0.00, 6.64	1.1 %		1/72	0/74	Lopaciuk 1992
0.36 [0.05, 2.61	4.3 %		3/85	1/85	Prandoni 1992
0.75 [0.17, 3.32	7.5 %		4/360	3/360	Prandoni 2004
0.51 [0.18, 1.40	16.1 %	•	715	721	Subtotal (95% CI)
			0.0%	$= 3 (P = 0.84); I^2 = (P = 0.19)$	Total events: 5 (LMWH), 10 (U Heterogeneity: $Chi^2 = 0.83$, df Test for overall effect: $Z = 1.31$
0.19 [0.06, 0.59	12.7 %		11/219	I/213	2 Tinzaparin versus unfractiona Hull 1992
0.61 [0.15, 2.46	8.6 %		5/308	3/304	Simonneau 1997
0.30 [0.12, 0.73	21.2 %	•	527	$= 1 (P = 0.20); I^2 = (P = 0.0078)$	Subtotal (95% CI) Total events: 4 (LMWH), 16 (U Heterogeneity: Chi ² = 1.67, df Test for overall effect: Z = 2.66 3 Enoxaparin versus unfraction
0.92 [0.33, 2.57	15.7 %	-	8/205	7/195	Decousus 1998
1.70 [0.42, 6.87	8.5 %	_	3/253	5/247	Levine 1996
Not estimab	0.5 /0		0/67	0/67	Simonneau 1993
1.14 [0.50, 2.61	24.2 %	_ _	525 0.0% 8/511	$= (P = 0.49); ^2 = (P = 0.76)$	Subtotal (95% CI) Total events: 12 (LMWH), 11 (Heterogeneity: Chi ² = 0.49, df Test for overall effect: Z = 0.31 4 Reviparin versus unfractionat Columbus 1997
_					
1.26 [0.49, 3.19	19.2 %	-	511	510 IFH)	Subtotal (95% CI) Total events: 10 (LMWH), 8 (U Heterogeneity: not applicable

(Continued ...)

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 128 thromboembolism (Review)

			Peto		(Continued)
Study or subgroup	LMWH	UFH	Odds Ratio	Weight	Peto Odds Ratio
	n/N	n/N	Peto,Fixed,95% Cl		Peto,Fixed,95% Cl
5 Certoparin versus unfraction	nated heparin				
Kirchmaier 1998	1/128	4/131		5.3 %	0.30 [0.05, 1.77]
Riess 2003	6/627	7/593		13.9 %	0.81 [0.27, 2.41]
Subtotal (95% CI)	755	724	•	19.3 %	0.62 [0.24, 1.56]
Total events: 7 (LMWH), 11 (U	JFH)				
Heterogeneity: $Chi^2 = 0.86$, df	$T = I (P = 0.35); I^2 =$:0.0%			
Test for overall effect: $Z = 1.02$	2 (P = 0.31)				
Total (95% CI)	3012	3002	•	100.0 %	0.68 [0.45, 1.03]
Total events: 38 (LMWH), 56 ((UFH)				
Heterogeneity: $Chi^2 = 10.64$, c	$df = 10 (P = 0.39); I^2$	2 =6%			
Test for overall effect: $Z = 1.84$	1 (P = 0.066)				
Test for subgroup differences: ($Chi^2 = 6.79, df = 4$	$(P = 0.15), ^2 = 41\%$			

0.001 0.01 0.1 1 10 100 1000 Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 129 thromboembolism (Review)

Analysis 8.6. Comparison 8 LMWH versus UFH: all randomised controlled trials with adequate concealment of allocation, Outcome 6 Overall mortality at the end of follow-up.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 8 LMWH versus UFH: all randomised controlled trials with adequate concealment of allocation

Outcome: 6 Overall mortality at the end of follow-up

Study or subgroup	LMWH	UFH	Peto Odds Ratio	Weight	Pet Odds Rati
	n/N	n/N	Peto,Fixed,95% CI		Peto,Fixed,95% (
I Nadroparin versus unfraction	ated heparin				
Koopman 1996	14/202	16/198	-	7.8 %	0.85 [0.40, 1.78
Lopaciuk 1992	0/74	1/72	•	0.3 %	0.13 [0.00, 6.64
Prandoni 1992	6/85	12/85		4.6 %	0.48 [0.18, 1.26
Prandoni 2004	12/360	12/360	+	6.5 %	1.00 [0.44, 2.26
Subtotal (95% CI)	721	715	•	19.2 %	0.76 [0.47, 1.22
Total events: 32 (LMWH), 41 (l Heterogeneity: Chi ² = 2.17, df Test for overall effect: Z = 1.13 2 Tinzaparin versus unfractiona	= 3 (P = 0.54); l ² = (P = 0.26)	=0.0%			
Hull 1992	10/213	21/219	-	8.1 %	0.48 [0.23, 1.00
Simonneau 1997	12/304	14/308	-	7.0 %	0.86 [0.39, 1.89
Subtotal (95% CI)	517	527	•	15.2 %	0.63 [0.37, 1.08
Heterogeneity: $Chi^2 = 1.15$, df Test for overall effect: $Z = 1.69$ 3 Ardeparin versus unfractional	(P = 0.091)	15/0			
3 Ardeparin versus unfractionat	` '				
Goldhaber 1998	0/41	0/39			Not estimab
Subtotal (95% CI) Total events: 0 (LMWH), 0 (UF Heterogeneity: not applicable Test for overall effect: not applie 4 Enoxaparin versus unfractiona	cable	39			Not estimabl
Decousus 1998	40/195	43/205	+	18.6 %	0.97 [0.60, 1.58
Levine 1996	11/247	17/253		7.5 %	0.65 [0.30, 1.40
Merli 2001	18/610	9/290	+	6.5 %	0.95 [0.42, 2.15
Simonneau 1993	3/67	2/67		1.4 %	1.51 [0.25, 8.96
Subtotal (95% CI) Total events: 72 (LMWH), 71 (I	1119 UFH) = 3 (P = 0.77); I ² :	815	•	33.9 %	0.90 [0.63, 1.29

Favours LMWH Favours UFH

(Continued . . .)

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 130 thromboembolism (Review)

Study or subgroup	LMWH n/N	UFH n/N	Peto Odds Ratio Peto,Fixed,95% Cl	Weight	(Continued) Peto Odds Ratio Peto,Fixed,95% Cl
Test for overall effect: $Z = 0.5$	6 (P = 0.57)				
5 Reviparin versus unfractiona	ted heparin				
Columbus 1997	36/510	39/511	•	19.6 %	0.92 [0.57, 1.47]
Subtotal (95% CI)	510	511	•	19.6 %	0.92 [0.57, 1.47]
Total events: 36 (LMWH), 39	(UFH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 0.3$	5 (P = 0.73)				
6 Certoparin versus unfraction	nated heparin				
Kirchmaier 1998	8/125	10/124		4.7 %	0.78 [0.30, 2.03]
Riess 2003	11/627	16/593		7.4 %	0.65 [0.30, 1.39]
Subtotal (95% CI)	752	717	•	12.2 %	0.70 [0.38, 1.26]
Total events: 19 (LMWH), 26	(UFH)				
Heterogeneity: $Chi^2 = 0.09$, d	$f = 1 (P = 0.76); I^2 =$	=0.0%			
Test for overall effect: $Z = 1.19$	9 (P = 0.23)				
Total (95% CI)	3660	3324	•	100.0 %	0.80 [0.65, 0.99]
Total events: 181 (LMWH), 21	2 (UFH)				
Heterogeneity: $Chi^2 = 6.32$, d	$f = 12 (P = 0.90); I^2$	=0.0%			
Test for overall effect: $Z = 2.0$	5 (P = 0.040)				
Test for subgroup differences:	$Chi^2 = 1.78, df = 4$	(P = 0.78), I ² =0.0%			

0.001 0.01 0.1 1 10 100 1000

Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 131 thromboembolism (Review)

Analysis 9.1. Comparison 9 LMWH versus UFH: all randomised controlled trials that used ISTH definition of major bleeding, Outcome 1 Incidence of major haemorrhagic episodes (during initial treatment).

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 9 LMWH versus UFH: all randomised controlled trials that used ISTH definition of major bleeding

Outcome: I Incidence of major haemorrhagic episodes (during initial treatment)

Pete Odds Ratio	Weight	Peto Odds Ratio	UFH	LMWH	Study or subgroup
Peto,Fixed,95% C		Peto,Fixed,95% Cl	n/N	n/N	
				nated heparin	I Nadroparin versus unfractio
Not estimable			0/196	0/98	Belcaro 1999
0.50 [0.05, 4.85	2.0 %		2/198	1/202	Koopman 1996
0.13 [0.00, 6.64	0.7 %	←	1/72	0/74	Lopaciuk 1992
0.48 [0.09, 2.43	3.8 %		4/81	2/85	Ninet 1991
0.36 [0.05, 2.61	2.6 %	<u> </u>	3/85	1/85	Prandoni 1992
0.75 [0.17, 3.32	4.6 %		4/360	3/360	Prandoni 2004
0.12 [0.01, 2.02	1.3 %		2/33	0/35	Thery 1992
0.44 [0.19, 1.01	14.9 %	•	1025	939 JFH)	Subtotal (95% CI) Total events: 7 (LMWH), 16 (1
			=0.0%	4 (P = 0.052)	Heterogeneity: Chi ² = 1.71, d Test for overall effect: Z = 1.9 2 Tinzaparin versus unfraction
0.19 [0.06, 0.59	7.7 %		11/219	1/213	Hull 1992
1.21 [0.60, 2.44	20.5 %	+	15/268	18/269	Leizorovicz 2011
0.61 [0.15, 2.46	5.2 %		5/308	3/304	Simonneau 1997
0.71 [0.41, 1.22	33.4 %	•	795	786	Subtotal (95% CI)
			=73%	$f = 2 (P = 0.02); I^2 = 4 (P = 0.21)$	Total events: 22 (LMWH), 31 Heterogeneity: Chi ² = 7.48, d Test for overall effect: Z = 1.2 3 Enoxaparin versus unfraction
0.92 [0.33, 2.57	9.5 %	-	8/205	7/195	Decousus 1998
Not estimable			0/30	0/29	Findik 2002
1.70 [0.42, 6.87	5.2 %		3/253	5/247	Levine 1996
5.61 [0.11, 297.44	0.6 %		0/21	1/29	P rez de Llano 2003
Not estimable			0/67	0/67	Simonneau 1993
1.22 [0.54, 2.75	15.4 %	•	576	567 (UFH)	Subtotal (95% CI) Total events: 13 (LMWH), 11

(Continued . . .)

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 132 thromboembolism (Review)

Study or subgroup	LMWH	UFH	Peto Odds Ratio	Weight	Peto Odds Ratio
	n/N	n/N	Peto,Fixed,95% Cl		Peto,Fixed,95% C
Heterogeneity: $Chi^2 = 1.08$, df		=0.0%			
Test for overall effect: $Z = 0.48$. ,				
4 Dalteparin versus unfraction		0.000		1.2.0/	
Fiessinger 1996	0/120	2/133		1.3 %	0.15 [0.01, 2.39
Lindmarker 1994	0/101	0/103			Not estimabl
Luomanmaki 1996	0/117	1/131	• · · · · · · · · · · · · · · · · · · ·	0.7 %	0.15 [0.00, 7.64
Meyer 1995	0/29	0/31			Not estimabl
Subtotal (95% CI)	367	398	-	2.0 %	0.15 [0.02, 1.44
Total events: 0 (LMWH), 3 (UI	FH)				
Heterogeneity: $Chi^2 = 0.00$, df	$I = I (P = 0.99); I^2$	=0.0%			
Test for overall effect: $Z = 1.6^{2}$. ,				
5 Reviparin versus unfractional					
Columbus 1997	10/510	8/511	-	11.7 %	1.26 [0.49, 3.19
Subtotal (95% CI)	510	511	+	11.7 %	1.26 [0.49, 3.19
Total events: 10 (LMWH), 8 (U	JFH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 0.48$	8 (P = 0.63)				
6 Certoparin versus unfraction	ated heparin				
Harenberg 2000a	4/265	11/273		9.6 %	0.40 [0.14, 1.10
Kirchmaier 1998	1/128	4/131		3.2 %	0.30 [0.05, 1.77
Riess 2003	6/627	7/593	-	8.5 %	0.81 [0.27, 2.41
Subtotal (95% CI)	1020	99 7	•	21.4 %	0.50 [0.25, 1.00
Total events: 11 (LMWH), 22 ((UFH)				
Heterogeneity: $Chi^2 = 1.26$, df	= 2 (P = 0.53); I ²	=0.0%			
Test for overall effect: $Z = 1.95$	5 (P = 0.052)				
7 Bemiparin versus unfractiona	ited heparin				
Kakkar 2003	2/111	0/110	+	1.3 %	7.39 [0.46, 8.89
Subtotal (95% CI)	111	110		1.3 %	7.39 [0.46, 118.89
Total events: 2 (LMWH), 0 (UI	FH)				
Heterogeneity: not applicable					
Test for overall effect: $Z = 1.41$	(P = 0.16)				
Total (95% CI)	4300	4412	•	100.0 %	0.71 [0.52, 0.98
Total events: 65 (LMWH), 91 (. ,				
Heterogeneity: Chi ² = 21.45, c	, , ,	12 = 16%			
Test for overall effect: $Z = 2.09$. ,				
Test for subgroup differences: (Chi² = 9.92, df = 6	$(P = 0.13), I^2 = 39\%$			

(... Continued)

Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 133 thromboembolism (Review)

Analysis 10.1. Comparison 10 LMWH versus UFH by year of publication, Outcome 1 Incidence of recurrent venous thromboembolism during initial treatment.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 10 LMWH versus UFH by year of publication

Outcome: I Incidence of recurrent venous thromboembolism during initial treatment

Study or subgroup	LMWH	UFH	Peto Odds Ratio	Weight	Pet Odds Rati
	n/N	n/N	Peto,Fixed,95% CI	-	Peto,Fixed,95% (
Ninet 1991	1/85	2/81		2.3 %	0.48 [0.05, 4.73
Prandoni 1992	1/85	4/85		3.9 %	0.29 [0.05, 1.72
Lopaciuk 1992	0/74	1/72	·	0.8 %	0.13 [0.00, 6.64
Thery 1992	0/35	0/33			Not estimab
Simonneau 1993	0/67	2/67		1.6 %	0.13 [0.01, 2.15
Lindmarker 1994	0/101	0/103			Not estimab
Meyer 1995	0/29	0/31			Not estimab
Levine 1996	7/247	12/253		14.5 %	0.59 [0.24, 1.48
Fiessinger 1996	1/111	1/120	<u> </u>	1.6 %	1.08 [0.07, 17.43
Koopman 1996	4/202	5/198	-	7.0 %	0.78 [0.21, 2.92
Simonneau 1997	3/304	2/308	<u> </u>	3.9 %	1.52 [0.26, 8.80
Columbus 1997	16/510	15/511	+	23.8 %	1.07 [0.52, 2.19
Kirchmaier 1998	1/128	3/131	<u> </u>	3.1 %	0.37 [0.05, 2.67
Decousus 1998	3/195	8/205		8.5 %	0.41 [0.13, 1.37
Harenberg 2000a	0/265	3/273		2.4 %	0.14[0.01, 1.34
Findik 2002	0/29	1/30	· · · · · · · · · · · · · · · · · · ·	0.8 %	0.14 [0.00, 7.06
Riess 2003	14/627	15/593	+	22.4 %	0.88 [0.42, 1.84
P rez de Llano 2003	3/29	2/21		3.5 %	1.09 [0.17, 6.98
otal (95% CI)	3123	3115	•	100.0 %	0.69 [0.49, 0.98
tal events: 54 (LMWH), 76 (UF eterogeneity: $Chi^2 = 9.77$, df = st for overall effect: $Z = 2.05$ (P est for subgroup differences: Not	4 (P = 0.78); ² = 0.04)				

0.001 0.01 0.1 1 10 100 1000 Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 134 thromboembolism (Review)

Analysis 10.2. Comparison 10 LMWH versus UFH by year of publication, Outcome 2 Incidence of recurrent venous thromboembolism at the end of follow-up.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 10 LMWH versus UFH by year of publication

Outcome: 2 Incidence of recurrent venous thromboembolism at the end of follow-up

Pe Odds Rat Peto,Fixed,95%	Weight	Peto Odds Ratio Peto,Fixed,95% Cl	UFH n/N	LMWH n/N	Study or subgroup
0.48 [0.18, 1.26	4.3 %		12/85	6/85	Prandoni 1992
0.42 [0.17, 1.0]	5.3 %		15/219	6/213	Hull 1992
0.13 [0.01, 1.25	0.8 %		3/72	0/74	Lopaciuk 1992
0.13 [0.01, 1.28	0.8 %		3/67	0/67	Simonneau 1993
1.71 [0.42, 7.0	2.0 %		3/103	5/101	Lindmarker 1994
Not estimat			0/31	0/29	Meyer 1995
0.79 [0.38, 1.6	7.5 %	-	17/198	14/202	Koopman 1996
0.77 [0.37, 1.6	7.4 %	-	17/253	13/247	Levine 1996
0.84 [0.26, 2.7	2.8 %		6/308	5/304	Simonneau 1997
1.09 [0.62, 1.9	13.0 %	+	25/511	27/510	Columbus 1997
0.87 [0.37, 2.0	5.5 %		12/205	10/195	Decousus 1998
0.13 [0.00, 6.4	0.3 %	•	1/39	0/41	Goldhaber 1998
0.50 [0.10, 2.5	1.5 %		4/124	2/125	Kirchmaier 1998
0.92 [0.34, 2.4	4.2 %		3/196	6/98	Belcaro 1999
0.42 [0.18, 1.0	5.3 %	_	15/273	6/265	Harenberg 2000a
-	7.1 %	_	13/2/3	21/610	Merli 2001
0.90 [0.42, 1.9					
0.31 [0.15, 0.6	7.8 %		24/375	7/388	Breddin 2001
0.36 [0.05, 2.7	1.0 %		3/30	1/29	Findik 2002
1.09 [0.17, 6.9	1.2 %		2/21	3/29	P rez de Llano 2003
0.76 [0.43, 1.3	12.4 %	1	27/593	22/627	Riess 2003
0.93 [0.44, 1.9	7.3 %	T	15/360	14/360	Prandoni 2004
2.25 [0.64, 7.8	2.6 %		3/268	7/269	Leizorovicz 2011
0.72 [0.59, 0.8	100.0 %	•	4621 ² = 13%	$f = 20 (P = 0.29); I^{2}$ (P = 0.0014)	btal (95% CI) tal events: 175 (LMWH), 23 eterogeneity: Chi ² = 23.04, c est for overall effect: $Z = 3.20$ est for subgroup differences: 1

Favours LMWH Favours UFH

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 135 thromboembolism (Review)

Analysis 10.3. Comparison 10 LMWH versus UFH by year of publication, Outcome 3 Incidence of major haemorrhagic episodes (during initial treatment).

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism Comparison: 10 LMWH versus UFH by year of publication

Outcome: 3 Incidence of major haemorrhagic episodes (during initial treatment)

Study or subgroup	LMWH	UFH	Peto Odds Ratio	Weight	Peto Odds Ratio
	n/N	n/N	Peto,Fixed,95% Cl		Peto,Fixed,95% Cl
Faivre 1988	0/33	3/35		1.9 %	0.13[0.01, 1.34]
Ninet 1991	2/85	4/81	_	3.8 %	0.48 [0.09, 2.43]
Lopaciuk 1992	0/74	1/72	·	0.6 %	0.13 [0.00, 6.64]
Hull 1992	1/213	11/219		7.6 %	0.19 [0.06, 0.59]
Thery 1992	0/35	2/33		1.3 %	0.12 [0.01, 2.02]
Prandoni 1992	1/85	3/85		2.5 %	0.36 [0.05, 2.61]
Simonneau 1993	0/67	0/67			Not estimable
Lindmarker 1994	0/101	0/103			Not estimable
Meyer 1995	0/29	0/31			Not estimable
Luomanmaki 1996	0/117	1/131	<	0.6 %	0.15 [0.00, 7.64]
Levine 1996	5/247	3/253		5.1 %	1.70 [0.42, 6.87]
Fiessinger 1996	0/120	2/133		1.3 %	0.15 [0.01, 2.39]
Koopman 1996	1/202	2/198		1.9 %	0.50 [0.05, 4.85]
Simonneau 1997	3/304	5/308		5.1 %	0.61 [0.15, 2.46]
Columbus 1997	10/510	8/511	+	11.4 %	1.26 [0.49, 3.19]
Kirchmaier 1998	1/128	4/131	<u> </u>	3.2 %	0.30 [0.05, 1.77]
Decousus 1998	7/195	8/205	-	9.4 %	0.92 [0.33, 2.57]
Belcaro 1999	0/98	0/196			Not estimable
Harenberg 2000a	4/265	11/273		9.4 %	0.40 [0.14, 1.10]
Findik 2002	0/29	0/30			Not estimable
Kakkar 2003	2/111	0/110		1.3 %	7.39 [0.46, 8.89]
P rez de Llano 2003	1/29	0/31		0.6 %	7.92 [0.16, 399.84]
Riess 2003	6/627	7/593		8.3 %	0.81 [0.27, 2.41]

0.001 0.01 0.1 1 10 100 1000

Favours LMWH Favours UFH

(Continued ...)

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 136 thromboembolism (Review)

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Study or subgroup	LMWH n/N	UFH n/N	Peto Odds Ratio Peto,Fixed,95% Cl	Weight	(Continued) Peto Odds Ratio Peto,Fixed,95% Cl
Prandoni 2004	3/360	4/360		4.5 %	0.75 [0.17, 3.32]
Leizorovicz 2011	18/269	15/268	+	20.1 %	1.21 [0.60, 2.44]
Total (95% CI)	4333	4457	•	100.0 %	0.69 [0.51, 0.95]
Total events: 65 (LMWH), 94	4 (UFH)				
Heterogeneity: Chi ² = 23.84	, df = 19 (P = 0.20); I	2 =20%			
Test for overall effect: $Z = 2$.	29 (P = 0.022)				
Test for subgroup differences	: Not applicable				
			0.001 0.01 0.1 1 10 100 1000		

Favours LMWH Favours UFH

Analysis 10.4. Comparison 10 LMWH versus UFH by year of publication, Outcome 4 Overall mortality at the end of follow-up.

Review: Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous thromboembolism

Comparison: 10 LMWH versus UFH by year of publication

Outcome: 4 Overall mortality at the end of follow-up

Study or subgroup	LMWH	UFH	Peto Odds Ratio	Weight	Peto Odds Ratio
	n/N	n/N	Peto,Fixed,95% CI		Peto,Fixed,95% Cl
Hull 1992	10/213	21/219	-#-	6.3 %	0.48 [0.23, 1.00]
Prandoni 1992	6/85	12/85		3.6 %	0.48 [0.18, 1.26]
Lopaciuk 1992	0/74	1/72	·	0.2 %	0.13 [0.00, 6.64]
Thery 1992	1/35	1/33		0.4 %	0.94 [0.06, 15.40]
Simonneau 1993	3/67	2/67		1.1 %	1.51 [0.25, 8.96]
Lindmarker 1994	2/101	3/103		1.1 %	0.68 [0.12, 3.99]
Meyer 1995	0/29	0/31			Not estimable
Koopman 1996	14/202	16/198	-	6.1 %	0.85 [0.40, 1.78]
Luomanmaki 1996	1/110	4/116		1.1 %	0.3 [0.05, .82]
			0.0010.010.1101001000		

Favours LMWH Favours UFH

(Continued ...)

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 137 thromboembolism (Review)

			Peto		(Continued Peto
Study or subgroup	LMWH UFH	UFH	Odds Ratio	Weight	Odds Ratio
	n/N	n/N	Peto,Fixed,95% Cl		Peto,Fixed,95% Cl
Levine 1996	11/247	17/253		5.8 %	0.65 [0.30, 1.40]
Columbus 1997	36/510	39/511	+	15.3 %	0.92 [0.57, 1.47]
Simonneau 1997	12/304	14/308	+	5.5 %	0.86 [0.39, 1.89]
Decousus 1998	40/195	43/205	+	14.5 %	0.97 [0.60, 1.58]
Kirchmaier 1998	8/125	10/124		3.7 %	0.78 [0.30, 2.03]
Goldhaber 1998	0/41	0/39			Not estimable
Harenberg 2000a	6/265	15/273		4.4 %	0.42 [0.18, 1.01]
Breddin 2001	9/388	11/375	-	4.3 %	0.79 [0.32, 1.91]
Merli 2001	18/610	9/290	+	5.0 %	0.95 [0.42, 2.15]
Findik 2002	0/29	0/30			Not estimable
Riess 2003	11/627	16/593		5.8 %	0.65 [0.30, 1.39]
Kakkar 2003	2/89	2/85		0.9 %	0.95 [0.13, 6.90]
P rez de Llano 2003	1/29	0/21		0.2 %	5.61 [0.11, 297.44]
Prandoni 2004	12/360	12/360	+	5.1 %	1.00 [0.44, 2.26]
Leizorovicz 2011	31/269	17/268	+	9.6 %	1.89 [1.04, 3.41]
otal (95% CI)	5004	4659	•	100.0 %	0.84 [0.70, 1.01]
otal events: 234 (LMWH), 2	65 (UFH)				
eterogeneity: Chi ² = 18.23,	df = 20 (P = 0.57);	2 =0.0%			
est for overall effect: $Z = 1.8$	2 (P = 0.068)				
est for subgroup differences:	Not applicable				

0.001 0.01 0.1 1 10 100 1000 Favours LMWH Favours UFH

ITS LITIVE Favour

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 138 thromboembolism (Review)

APPENDICES

Appendix I. CENTRAL search strategy

#1	MESH DESCRIPTOR Thrombosis	1231
#2	MESH DESCRIPTOR Thromboembolism	892
#3	MESH DESCRIPTOR Venous Thromboembolism	233
#4	MESH DESCRIPTOR Venous Thrombosis EXPLODE ALL TREES	1996
#5	(thromboprophyla* or thrombus* or thrombotic* or throm- bolic* or thromboemboli* or thrombos* or embol*):TI,AB, KY	17001
#6	MESH DESCRIPTOR Pulmonary Embolism EXPLODE ALL TREES	729
#7	(PE or DVT or VTE):TI,AB,KY	4480
#8	(((vein* or ven*) near thromb*)):TI,AB,KY	6111
#9	#1 OR #2 OR #3 OR #4 OR #5 OR #6 OR #7 OR #8	20325
#10	MESH DESCRIPTOR Heparin EXPLODE ALL TREES	3815
#11	heparin*:TI,AB,KY	8661
#12	LMWH:TI,AB,KY	790
#13	UFH:TI,AB,KY	437
#14	UH:TI,AB,KY	84
#15	(nadroparin* or fraxiparin* or enoxaparin or Clexane or klex- ane or lovenox or dalteparin or Fragmin or ardeparin or normi- flo or tinzaparin or logiparin or Innohep or certoparin or san- doparin or reviparin or clivarin* or danaproid or danaparoid) :TI,AB,KY	2405
#16	(antixarin or ardeparin* or bemiparin* or Zibor or cy 222 or embolex or monoembolex or parnaparin* or "rd 11885" or tedelparin or Kabi-2165 or Kabi 2165):TI,AB,KY	149
#17	(emt-966 or emt-967 or "pk-10 169" or pk-10169 or pk10169):TI,AB,KY	8

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(Continued)

#18	(fr-860 or cy-216 or cy216 or seleparin* or tedegliparin or seleparin* or tedegliparin*):TI,AB,KY	51
#19	("kb 101" or kb101 or lomoparan or orgaran):TI,AB,KY	31
#20	(parnaparin or fluxum or lohepa or lowhepa or "op 2123" or parvoparin or AVE5026):TI,AB,KY	36
#21	#10 OR #11 OR #12 OR #13 OR #14 OR #15 OR #16 OR #17 OR #19 OR #20	9580
#22	#9 AND #21	4334

Appendix 2. Trials registries searches

Clinicaltrials.gov 134 studies found for: subcutaneous AND heparin WHO 57 records for 42 trials found for: subcutaneous AND heparin ISRCTN 13 results subcutaneous AND heparin

FEEDBACK

Anticoagulant feedback, 14 February 2011

Summary

Feedback received on this review, and other reviews and protocols on anticoagulants, is available on the Cochrane Editorial Unit website at http://www.editorial-unit.cochrane.org/anticoagulants-feedback.

WHAT'S NEW

Last assessed as up-to-date: 15 September 2016.

Date	Event	Description
15 September 2016	New search has been performed	Searches rerun. Six new studies included, five new studies excluded and one ongoing study identified
15 September 2016	New citation required but conclusions have not changed	Searches rerun. Six new studies included, five new studies excluded and one ongoing study identified. Review updated according to current Cochrane stan-

Fixed dose subcutaneous low molecular weight heparins versus adjusted dose unfractionated heparin for the initial treatment of venous 140 thromboembolism (Review)

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dards. New authors have taken over this review. Conclusions not changed

HISTORY

Protocol first published: Issue 3, 1997 Review first published: Issue 2, 1998

Date	Event	Description
14 February 2011	Amended	Link to anticoagulant feedback added
14 July 2010	New search has been performed	The review was updated, one additional trial was added to the included studies and two additional trials were excluded
27 April 2010	New citation required but conclusions have not changed	There was a change in authors in the updated review.
20 October 2008	Amended	Converted to new review format.
14 November 2005	Amended	Minor copy edits made.
23 August 2004	New citation required but conclusions have not changed	Change in authors.
23 August 2004	New search has been performed	Review substantively updated by the addition of eight new included studies. Conclusions unchanged
15 February 1999	New search has been performed	One additional trial included but no change to con- clusions.

CONTRIBUTIONS OF AUTHORS

LR: selected studies for inclusion in this update, assessed the quality of studies, carried out data extraction, performed data analysis and wrote the review.

LJ: selected studies for inclusion in this update, assessed the quality of the studies and carried out data extraction.

DECLARATIONS OF INTEREST

LR: none known.

LJ: none known.

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• No sources of support supplied

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• Chief Scientist Office, Scottish Government Health Directorates, The Scottish Government, UK.

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DIFFERENCES BETWEEN PROTOCOL AND REVIEW

Included post hoc sensitivity analysis for ISTH bleeding definitions in order to assess the effect of bleeding definitions used.

INDEX TERMS

Medical Subject Headings (MeSH)

Anticoagulants [*administration & dosage; adverse effects]; Hemorrhage [chemically induced]; Heparin [administration & dosage; adverse effects]; Heparin, Low-Molecular-Weight [*administration & dosage; adverse effects]; Injections, Subcutaneous; Pulmonary Embolism [*drug therapy; mortality]; Randomized Controlled Trials as Topic; Recurrence; Venous Thrombosis [*drug therapy; mortality]

MeSH check words

Humans