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Randomized trials of therapeutic heparin for COVID-19: A meta-analysis

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Essentials

- Trials suggest differences in the effect of therapeutic heparin based on severity of COVID-19.
- We did a meta-analysis to determine the effect of therapeutic heparin in hospitalized patients.
- In the moderately ill, there was a significant reduction in death or mechanical ventilation.
- In the severely ill, there was no evidence for benefit of therapeutic heparin.

1 | INTRODUCTION

Pulmonary inflammation, endothelial injury, and microcirculatory thromboses likely contribute to hypoxemic respiratory failure, the most common cause of death in patients with COVID-19. Observational data suggest that heparin anticoagulation decreases the risk of critical illness and death in those hospitalized for COVID-19.\(^1\)\(^2\) Heparin anticoagulants are of particular interest for the treatment of COVID-19 due to their additional anti-inflammatory and potentially antiviral properties.\(^6\)\(^-\)\(^8\) Randomized controlled trials (RCTs) suggest that therapeutic heparin anticoagulation is beneficial in patients hospitalized for COVID-19 with moderate illness, but of no benefit and potential harm when provided to patients with critical illness.\(^9\)\(^10\) Given the disparate findings in these two patient populations and safety concerns regarding bleeding, there is hesitancy to adopt therapeutic heparin as standard care in moderately...
ill patients with COVID-19.\textsuperscript{11} We conducted a systematic review and meta-analysis of available RCTs of therapeutic-dose heparin anticoagulation compared with usual care to determine the effects of therapeutic heparin in hospitalized patients with COVID-19.

2 | METHODS

This systematic review and meta-analysis was reported according to the Preferred Reporting for Systematic Reviews and Meta-Analyses guidelines.\textsuperscript{12} We searched PubMed, Embase, and Web of Science, using terms (“heparin” OR “dalteparin” OR “enoxaparin” OR “tinzaparin” OR “anticoagulation”) AND (“SARS-CoV-2” OR “COVID” OR “coronavirus” OR “COVID-19” AND (“randomized” OR “randomised” OR “clinical trials”), with no language restrictions. We also searched the grey literature, which included medRxiv, and medical conference proceedings. We included RCTs comparing therapeutic-dose heparin anticoagulation with usual care in hospitalized adults with moderate or severe COVID-19. Moderate illness was defined as admission to hospital ward level of care, not already mechanically ventilated, and not imminently requiring mechanical ventilation or critical care. Severe illness was defined as admission to hospital with clinically intensive level of care. We restricted the search to RCTs published from March 1, 2019, to October 8, 2021. We excluded trials that used oral anticoagulation or used intermediate dosing of heparin in the experimental arm. Heparin was defined as either unfractionated intravenous or low-molecular-weight subcutaneous forms of heparin; both forms are within the same drug class, exert their anticoagulant effect by amplifying the activity of antithrombin and have similar nonanticoagulant effects.\textsuperscript{13} Two reviewers (MS and GT) independently screened title, abstract, and full text of retrieved articles for inclusion. Any disagreement or uncertainty was resolved by consensus. We used the Cochrane risk of bias tool to assess risk of bias in the included trials.\textsuperscript{14}

Prespecified outcomes included all-cause death, death or invasive mechanical ventilation, death or organ support, death or major thrombotic event, death or any thrombotic event, major thrombotic events, major bleeding as defined by the ISTH,\textsuperscript{15} ventilator-free days alive, and organ support-free days alive. Major thrombotic events were defined as the composite of myocardial infarction, pulmonary embolism, ischemic stroke, or systemic arterial embolism. Any thrombotic events were defined as major thrombotic events or deep vein thromboses. Definition of organ support-free days alive are described in detail in the published protocols of the included clinical trials.\textsuperscript{16,17} Ventilator-free days, and organ support-free days alive were analyzed using ordinal logistic regression; death was assigned the worst outcome (a value of −1) in these analyses.\textsuperscript{18}

We used forest plots to display the results of the meta-analysis. Mantel-Haenszel fixed-effect meta-analyses were used to combine odds ratios (ORs) of outcomes reported in available RCTs separately for moderately ill ward patients and severely ill intensive care unit (ICU) patients, using a chi-squared test to estimate P values for interaction between treatment and severity of illness. The variance attributed to pooled results reflects only sampling error due to the play of chance at randomization. Homogeneity of ORs is not required for fixed-effect pooled ORs to be informative.\textsuperscript{19} Heterogeneity was evaluated using I-square values. I-square values of >25% suggests low, 50% moderate, and 75% high between-trial heterogeneity.\textsuperscript{20} Statistical analyses were performed using Stata version 15 (StataCorp, College Station, TX, USA).

3 | RESULTS AND DISCUSSION

There were four trials that compared therapeutic heparin to usual care using lower doses of heparin (see Figure 1).\textsuperscript{9,10,21,22} Three trials included moderately ill ward patients with COVID-19: a multiplatform trial integrating the Antithrombotic Therapy to Ameliorate Complications of COVID-19 (ATTACC), Accelerating COVID-19 Therapeutic Interventions and Vaccines-4 Antithrombotics Inpatient Platform Trial (ACTIV-4a) and the Randomized, Embedded, Multifactorial Adaptive Platform Trial for Community-Acquired Pneumonia (REMAP-CAP)\textsuperscript{9} the Coagulopathy of COVID-19, A Pragmatic Randomized Controlled Trial of Therapeutic Anticoagulation Versus Standard Care as a Rapid Response to the COVID-19 Pandemic (RAPID),\textsuperscript{21} and the Systemic Anticoagulation With Full Dose Low Molecular Weight Heparin (LMWH) Versus Prophylactic or Intermediate Dose LMWH in High Risk COVID-19 Patients (HEP-COVID) trial.\textsuperscript{22} Three trials included severely ill patients with COVID-19: a separate multiplatform trial conducted by the same investigators evaluated therapeutic heparin in severely ill ICU patients,\textsuperscript{23} a small phase II randomized trial of mechanically ventilated patients (Therapeutic Versus Prophylactic Anticoagulation for Severe COVID-19: A Randomized Phase II Clinical Trial [HESACOVID]),\textsuperscript{24} and the HEP-COVID trial, which also included severely ill patients, with randomization stratified according to disease severity.\textsuperscript{22} We excluded the Therapeutic versus Prophylactic Anticoagulation for Patients Admitted to Hospital with COVID-19 and Elevated D-dimer Concentration (ACTION) trial, as it combined therapeutic anticoagulation with rivaroxaban in moderately ill patients and therapeutic enoxaparin in severely ill patients, without reporting any of our prespecified outcomes by illness severity.\textsuperscript{25} We also excluded the Effect of Intermediate-Dose Versus Standard-Dose Prophylactic Anticoagulation on Thrombotic Events, Extracorporeal Membrane Oxygenation Treatment, or Mortality Among Patients With COVID-19 Admitted to the Intensive Care Unit (INSPIRATION)\textsuperscript{26} and the trial of Perepu et al.\textsuperscript{27} as these trials used intermediate dose heparin in their experimental arms. The risk of bias assessment of the included trials is presented in Table S1. There were deviations from the intended experimental intervention in the two multiplatform trials where lower-than-therapeutic doses of heparin were administered in >20% of patients allocated to the experimental arms.\textsuperscript{9,10} Additionally, the outcomes of organ support were not blindly adjudicated by an independent clinical events committee in the multiplatform trials. There was no information on protocol deviations, and no prespecified statistical analysis plan provided or information on blinded evaluation of the secondary outcomes in the HESACOVID trial.\textsuperscript{24} The risk of bias was considered low for the RAPID and the HEP-COVID trials.\textsuperscript{21,22}
Figure 2 shows meta-analyses of three trials in moderately ill patients. Data from the HEP-COVID trial contributed to the meta-analysis of the outcomes of death or any thrombotic event and major bleeding, as to date only these outcomes were presented by illness severity. There was nonsignificant reduction in all-cause death (OR, 0.76; 95% CI, 0.57-1.02), but significant reductions in the composite of death or invasive mechanical ventilation (OR, 0.77; 95% CI, 0.60-0.98), death or organ support (OR, 0.77; 95% CI, 0.63-0.93), death or major thrombotic event (OR, 0.64; 95% CI, 0.48-0.86), death or any thrombotic event (OR, 0.58; 95% CI, 0.45-0.77), and major thrombotic events (OR, 0.47; 95% CI, 0.24-0.90). Ventilator-free days alive (OR, 1.30; 95% CI 1.05-1.61) and organ support-free days alive (OR, 1.29; 95% CI, 1.07-1.57) were significantly increased with therapeutic heparin. Conversely, there was a nonsignificant increase in major bleeding. Figure 3 shows meta-analyses of the three trials in severely ill patients.

Figure 4 shows analyses of the interaction between treatment effect and severity of illness. In severely ill patients, there was no evidence to suggest a benefit of therapeutic heparin. There were significant treatment-by-subgroup interactions with severity of illness for all-cause death ($P$ for interaction = .034), all-cause death or major thrombotic event ($P = .017$), death or any thrombotic event ($P = 0.002$), and organ support-free days alive ($P = .003$). There was no evidence for treatment-by-subgroup interactions for major thrombotic events and major bleeding.

This meta-analysis included three available trials of therapeutic heparin in moderately ill ward patients with COVID-19, which were not individually conclusive. Findings for five effectiveness outcomes (death or invasive mechanical ventilation, death or organ support, ventilator-free days alive, organ support-free days alive, major thrombotic events) were consistent between trials, with significant differences in favor of therapeutic heparin. There was high heterogeneity in the findings for all-cause death and the composites of death or major thrombotic event, and death or any thrombotic event. This suggests that there are competing nonmodifiable causes of death aside from micro- and macrovascular thromboses. By contrast to positive treatment effects in ward patients, there were no such effects in severely ill ICU patients. We found significant treatment-by-subgroup interactions with severity of illness for all-cause death, all-cause death or major thrombosis, all-cause death or any thrombosis, and organ support-free days alive, with evidence of benefit with therapeutic heparin in moderately ill ward patients, but not in severely ill ICU patients. Conversely, there were no treatment-by-subgroup interactions for major thrombotic events and major bleeding, with benefit in both groups for therapeutic heparin to prevent major thrombotic events with non-significant,
FIGURE 2 Meta-analyses of effectiveness and safety outcomes in randomised trials comparing therapeutic heparin with usual care in moderately ill ward patients with COVID-19. Mantel-Haenszel fixed-effect meta-analyses of the RAPID trial, the multiplatform trial and HEP-COVID trial in moderately ill ward patients.9,21,22 Squares and horizontal lines show treatment effects and their 95% confidence intervals in each trial. The area of each square is proportional to the weight the trial received in the meta-analysis. Diamonds show estimated treatment effects and 95% confidence intervals from meta-analyses. Odds ratios for ventilator-free and organ support–free days alive are from ordinal logistic regression in both trials; death was assigned the worst outcome (a value of −1). Absolute values were not available for all-cause death from the multiplatform trial. Major thrombotic events were defined as the composite of myocardial infarction, pulmonary embolism, ischemic stroke or systemic arterial embolism; any thrombotic events were defined as a major thrombotic event or deep vein thrombosis; major bleeding defined by the ISTH Scientific and Standardization Committee. The observation time for the outcomes in the trials were 28 days for the multiplatform trial (with the exception of organ support–free days, which was calculated for an observation time of 21 days), 28 days for the RAPID trial, and 30 days for HEP-COVID trial.6,17,23
numeric increases in major bleeding. We speculate that the different effect of therapeutic heparin in moderately ill compared to severely ill patients was because the latter were too ill for this treatment to alter the cascade of endothelial injury and microvascular thromboinflammation, emphasizing the need for early treatment. However, there were consistent and significant reductions in major thrombotic events in the severely and moderately ill, which suggests that therapeutic heparin is an effective anticoagulant protecting against large-vessel thrombosis and thromboembolism in patients with COVID-19.28,29

A recently published meta-analysis by Ortega-Paz et al20 differs from ours, concluding that prophylactic anticoagulation should be preferred over intermediate or therapeutic dose anticoagulation for hospitalized patients with COVID-19, whether moderately or severely ill. The authors combined trials using different types and doses of anticoagulants in the experimental arm, including

FIGURE 3 Meta-analyses of effectiveness and safety outcomes in randomised trials comparing therapeutic heparin with usual care in severely ill intensive care unit (ICU) patients with COVID-19. Mantel-Haenszel fixed-effect meta-analyses of the multiplatform trial, HESACOVID and HEP-COVID in severely ill ICU patients.10,22,24 Squares and horizontal lines show treatment effects and their 95% confidence intervals in each trial. The area of each square is proportional to the weight the trial received in the meta-analysis. Diamonds show estimated treatment effects and 95% confidence intervals from meta-analyses. Odds ratios for organ support–free days alive are from ordinal logistic regression; death was assigned the worst outcome (a value of −1). Major thrombotic events were defined as the composite of myocardial infarction, pulmonary embolism, ischemic stroke, or systemic arterial embolism; any thrombotic events were defined as a major thrombotic event or deep vein thrombosis; major bleeding defined by the ISTH Scientific and Standardization Committee. The observation time for the outcomes in the trials were 28 days for the multiplatform trial (with the exception of organ support–free days, which was calculated for an observation time of 21 days), and 30 days for HEP-COVID trial10,23
rivaroxaban and intermediate- or therapeutic-dose heparin. This variation in type and dose of anticoagulant is likely associated with variation in treatment effect, as different types and doses of anticoagulants have different mechanisms of action, off-target effects, and safety profiles. Pooling trials with different experimental regimens could therefore submerge important treatment effects of single regimens. Our conclusions differ, as we only pooled data from trials of therapeutic- compared to lower dose-dose heparin. Also, Ortega-Paz et al evaluated effectiveness only for the outcomes of all-cause death and large-vessel thrombosis, while we also analyzed outcomes such as organ support-free days alive and the composite of death or thrombosis, which more completely evaluate the effect of pulmonary and systemic thromboinflammation on health and health resource use.

A limitation of this meta-analysis, like the majority of published meta-analyses, is that it did not analyze patient-level data. Only five
trials were included, which may be considered a limitation, but early and timely synthesis of trial-level data is important in the setting of the ongoing COVID-19 pandemic. A strength of this research is the harmonized definitions used in three of the included trials.\textsuperscript{9,21,22} Also, the fact that the results of these different multicenter trials (conducted in different settings) are generally in agreement further supports the robustness of the findings of this meta-analysis. There are multiple other ongoing trials evaluating therapeutic versus prophylactic anticoagulation, so an updated meta-analysis will be important.\textsuperscript{31,32} However, in the face of the ongoing pandemic, our findings provide sufficient evidence to support the adoption of therapeutic heparin as standard care in moderately ill hospitalized patients with COVID-19 who are at low risk of bleeding. In conclusion, this meta-analysis showed that therapeutic heparin is beneficial in moderately ill ward patients, but not in severely ill patients hospitalized with COVID-19.

**RELATIONSHIP DISCLOSURE**

The authors declare no conflicts of interest.

**AUTHOR CONTRIBUTIONS**

Concept and design: MS, BRdC, MC, PJ. Acquisition, analysis, or interpretation of data: MS, BRdC, PJ. Drafting of the manuscript: MS, BRdC, M Cushman, PJ. Critical revision of the manuscript: MS, BRdC, GHT, HR, MAH, LBK, FNÁ, MOA, PDJ, DL, M Carrier, AB, MF, SM, AYYL, KET, EMN, M Cushman, PJ. Statistical analysis: BRdC. All authors approved the final manuscript and submission.

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**SUPPORTING INFORMATION**

Additional supporting information may be found in the online version of the article at the publisher's website.

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