COVID-19 patient plasma demonstrates resistance to tPA-induced fibrinolysis as measured by thromboelastography

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Abstract
Patients critically ill with COVID-19 are at risk for thrombotic events despite prophylactic anticoagulation. Impaired fibrinolysis has been proposed as an underlying mechanism. Our objective was to determine if fibrinolysis stimulated by tissue plasminogen activator (tPA) differed between COVID patients and controls. Plasma from 14 COVID patients on prophylactic heparin therapy was obtained and compared with heparinized plasma from 14 different healthy donors to act as controls. Kaolin activated thromboelastography with heparinase was utilized to obtain baseline measurements and then repeated with the addition of 4 nM tPA. Baseline fibrinogen levels were higher in COVID plasma as measured by maximum clot amplitude (43.6 ± 6.9 mm vs. 23.2 ± 5.5 mm, p < 0.0001) and Clauss assay (595 ± 135 mg/dL vs. 278 ± 44 mg/dL, p < 0.0001). With the addition of tPA, fibrinolysis at 30 min after MA (LY30%) was lower (37.9 ± 16.5% vs. 58.9 ± 18.3%, p = 0.0035) and time to 50% lysis was longer (48.8 ± 16.3 vs. 30.5 ± 15.4 min, p = 0.0053) in the COVID-19 samples. Clotting times and rate of fibrin polymerization (‘R’ or ‘α’ parameters) were largely the same in both groups. Clot from COVID patients contains a higher fibrin content compared to standard controls and shows resistance to fibrinolysis induced by tPA. These findings suggest the clinical efficacy of thrombolytics may be reduced in COVID-19 patients.

Keywords Thromboelastography · Fibrinolysis · COVID-19 · Severe acute respiratory syndrome coronavirus 2 · Tissue plasminogen activator

Introduction
Patients with COVID-19, the disease caused by the severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), are at increased risk for venous thrombotic events (VTE) despite prophylactic anticoagulation [1]. The hypercoagulability profile is characterized by very high fibrinogen and d-dimer levels, elevated Factor VIII and von Willebrand factor levels, as well as reduced measurements of natural anticoagulants such as Protein C, Protein S, and antithrombin [2]. Viscoelastic testing (i.e. thromboelastography (TEG) or rotational thromboelastometry) typically demonstrates increased clot strength with a higher fibrin contribution [3–5].

Perhaps somewhat paradoxically in light of their elevated d-dimers, COVID-19 patients are also thought to exhibit reduced fibrinolytic activity. “Fibrinolytic shutdown” as defined by viscoelastic testing has been reported in these patients and is associated with an increased incidence of VTE [6, 7]. It has been suggested that fibrinolytic agents may be a therapeutic option to treat this impaired fibrinolysis
Micro clots in the lungs may also be important in the development of sepsis-related acute respiratory distress syndrome (ARDS). In a phase I clinical trial in trauma patients, Hardaway et al. demonstrated improved PaO2 with streptokinase administration in patients suffering from ARDS, presumably from dissolution of these micro clots [8]. This reasoning formed the basis for a recent small case series of three intubated COVID-19 patients with ARDS treated with low dose tissue plasminogen activator (tPA) in which improvement of P/F ratios were observed [9].

Given the increased clot strength described in COVID-19, it is possible that the effects of fibrinolytic therapy in these patients may be reduced. We aimed to compare the fibrinolytic response of tPA using TEG measurements on patient and control plasma. We hypothesized that COVID-19 patients would show reduced clot lysis percentage compared to controls.

Materials and methods

Sample collection

Platelet poor plasma (PPP) was obtained from patients diagnosed with acute COVID-19 and admitted to Emory University Hospital between 6/28/2020 and 7/11/2020. All samples were collected within 8 days of admission. Any associated clinical labs were collected in a deidentified fashion according to a protocol approved by Emory’s Institutional Review Board. All patients were on prophylactic anticoagulation with either low molecular weight or unfractionated heparin at the time of collection. Patients on any other anticoagulation medications were excluded. Plasma from 14 different healthy individuals (7 male and 7 female) was purchased (George King Bio-Medical Inc., Overland Park, KS) to act as controls. Prior to testing, unfractionated heparin (1.5 units/mL) was added to each control sample to replicate patient anticoagulation. All samples were stored at −80 °C until thawed for testing.

Thromboelastography

All testing was performed on the TEG 5000 (Haemonetics, Boston, MA) using heparinase cups containing 2.0 IU lyophilized Heparinase I, which is capable of neutralizing heparin levels of up to 6.0 u/mL according to the manufacturer. This has been previously shown to neutralize both unfractionated and low molecular weight heparins (LMWH) for TEG testing [10]. Patient and control plasma were tested both with and without the addition of tPA (final concentration 4 nM, Activase alteplase, Genentech, San Francisco, CA) to the cup.

Our primary measurements of interest were those TEG parameters related to fibrinolysis. The Ly30 and Ly60 parameters represent the percentage decrease in amplitude that occurs 30 and 60 min respectively after the maximum amplitude (MA) is reached [11]. The higher the percentage, the greater amount of fibrinolysis has taken place. The MA, measured in millimeters, represents clot strength and is normally a function of platelet count and fibrinogen interactions, with higher levels of each associated with greater clot amplitude. However, when PPP is used, the clot amplitude is dependent only on fibrinogen interactions. The clot lysis time (CLT) is another parameter of fibrinolysis and represents the elapsed time from when MA is reached to 50% clot lysis. Although not related to fibrinolysis, we collected the R time (R), which is a clotting time that begins with the addition of calcium and kaolin to the citrated sample and ends with initial fibrin formation. The alpha angle (α), which is a function of fibrin polymerization rate, was also recorded.

Statistical analysis

A convenience sample of COVID-19 patients was planned. Normality of the data was assessed using the Kolmogorov–Smirnov test and visual inspection of Q-Q plots. Unpaired t tests or Wilcoxon’s exact test as appropriate were used to compare continuous variables between groups. Probability values of α < 0.05 were required to reject the null hypothesis in a two-tailed test. All calculations were made using SAS 9.4 (SAS Institute, Cary, NC) and Prism 9.0 (GraphPad Software, San Diego, CA).

Results

Samples were collected from 14 different patients, 8 of whom were admitted to the ICU and 6 of whom were on the hospital ward floor. All samples were collected within 8 days of admission (range 1–8). Floor patients were exclusively on LMWH anticoagulation while the majority of ICU patients were on infusions of unfractionated heparin. Available laboratory results for these patients are provided in Table 1. Fibrinogen by Clauss assay was also measured in the 14 samples from healthy controls. Samples from COVID-19 patients contained approximately twice as much fibrinogen (594 ± 135 vs. 278 ± 44 mg/dL, p < 0.001).

The TEG measurements with and without the addition of tPA are presented in Table 2. Like the Clauss fibrinogen level, the MA parameter was also twice as high in COVID-19 samples. The R was also longer in the COVID-19 plasma. Neither COVID-19 nor control plasma showed any fibrinolysis at 30 min in the standard heparinase cups and thus LY60 and CLT measurements were not obtained.
Both groups showed a marked increase in LY30 and Ly60 with the addition of tPA. The mean amount of clot lysis in COVID-19 patients was 21% less than controls at 30 min and 15% less at 60 min. The mean CLT was more than 50% longer in COVID-19 samples compared to controls. All fibrinolysis measurements are presented graphically in Fig. 1. For other TEG parameters, only the MA was significantly higher in the COVID-19 patients.

COVID-19 patients are known to display a hypercoagulable profile on whole blood TEG consistent with increased clot strength and high fibrinogen levels [12]. Our sample of patients included ICU and hospital ward floor who, by C-reactive protein measurements, would be classified as having at least moderate or severe disease [13]. Since our plasma samples contained no platelets, the measured clot strength in this study was entirely dependent on fibrin interactions. The higher MA measured in COVID-19 samples both with and without tPA addition was expected. This was also confirmed with the baseline differences in fibrinogen measurements by Clauss assay, which is consistent with previously reported hematological findings of this disease.

Our primary aim was not to demonstrate increased fibrinogen levels, but to investigate the effects of tPA on fibrinolysis in COVID-19 patients. However, the optimal dose to use for this purpose in plasma-based TEG is unclear. In the trauma literature, Moore et al. have utilized doses of 75 ng/mL (~1.1 nM) for “tPA challenge TEG” in whole blood, which would be equivalent to approximately 2 nM in plasma [14]. Ostrowski et al. utilized 1.8 nM tPA when comparing plasma from patients with the human immune-deficiency virus (HIV) to controls [15]. They obtained Ly30 values of 24.2% in healthy controls (vs. 53.9% in hypocoagulable HIV samples) [15]. Given the hypercoagulable profile of COVID-19, we were not sure if these previously reported doses would show any effect, so we decided to use 4 nM tPA, which was about twice the concentration of these prior studies. Like Ostrowski et al., we saw no fibrinolysis on the standard TEG in both groups with and without tPA addition was expected. This was also confirmed with the baseline differences in fibrinogen measurements by Clauss assay, which is consistent with previously reported hematological findings of this disease.

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Decreased fibrinolysis in COVID-19 patients was also recently reported by Bachler et al. using whole blood on
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The Clot Pro (Enicor GmbH, Munich, Germany) viscoelastic platform [16]. They reported a more than doubled lysis time (LT) in COVID-19 patients compared to healthy controls. The LT parameter is very similar to the TEG CLT parameter that we measured, although the dose of tPA in the Clot Pro is approximately 2.5 times higher than what was used in our study. While it is difficult to translate values across different viscoelastic testing platforms, it is notable that our study in plasma demonstrated about the same magnitude of resistance to fibrinolysis as Bachler et al.

There are several possibilities for the reduced fibrinolysis demonstrated in the COVID-19 samples. One plausible explanation is that there are increased levels of fibrinolytic inhibitors present. Plasminogen activator inhibitor-1 (PAI-1), for example, has been shown to be elevated in these patients [17]. The mechanism is thought to be due to the downregulation of angiotensin converting enzyme by the coronavirus (SARS-CoV-2), leading to increased levels of angiotensin II, an inducer of PAI-1 in endothelial cells [18].

Another possibility may simply be the high fibrinogen levels lead to clot formation with a dense fibrin structure, making it less susceptible to tPA-induced fibrinolysis [19]. Increased fibrinogen in COVID-19 patients has been shown to correlate with levels of the inflammatory cytokine IL-6, which is also associated with more severe disease [20, 21]. Recently, our group reported on therapeutic plasma exchange in six critically ill COVID-19 patients which reduced their fibrinogen levels by about 50% [22]. Notably, the fibrinogen levels remained low and four of the six patients had clinical improvement. It is possible that simply reducing fibrinogen levels may decrease the sequelae of COVID-19 hypercoagulability. By decreasing clot strength within the microvasculature, the natural actions of the fibrinolytic system may be more effective.

Despite the high fibrinogen levels, our study did not demonstrate significantly different values for the α angle between COVID-19 and control plasma. This perhaps suggests that the inhibition of fibrinolysis may be more important than...
an increased rate of fibrin formation in the development of COVID-19 hypercoagulability. Although there was a difference in R between COVID-19 and control plasma at baseline, this was not reproduced in the TEGs with tPA addition. In general, COVID-19 patients are not thought to have exceptionally prolonged clotting times [5], and it is possible the prolonged R observed on baseline measurements was simply due to chance or perhaps less effective heparin neutralization in that particular batch of TEG cups.

This was a small ex-vivo study with some obvious limitations. Other than being in the ICU or on the wards, we were not able to collect much demographic information given that the samples were deidentified. It is possible that other comorbidities in addition to COVID-19 existed in these patients that could have increased their thrombogenicity and resistance to clot lysis. Additionally, since it was performed in platelet-poor plasma, the TEG measurements only reflect fibrinogen contributions to clot strength, ignoring the role of platelets. Some studies have suggested that platelets can enhance clot resistance to fibrinolysis [23]. Prior whole blood viscoelastic studies in COVID-19 patients have shown an increased platelet contribution to clot strength, but this still seems to be less prominent than the fibrinogen contribution [20]. Some authors have reported increased aggregation in COVID-19 platelets possibly due to increased thromboxane generation [24]. However, aspirin does not appear to prevent COVID-19 thrombosis [25]. Further studies are warranted to investigate the role of platelets in COVID-19 hypercoagulability.

In conclusion, we have demonstrated an increased resistance to tPA-induced fibrinolysis in COVID-19 plasma. This may be due, at least in part, to simply having a higher fibrinogen content, although other mechanisms are possible. The reduced efficacy of tPA may have clinical implications. As described in a recent case report, COVID-19 clot dissolution can require more than 100 mg of tPA and catheter-directed treatment as well [26]. Typical dosing schemes of fibrinolytics may need to be increased, but further research is required.

Author contributions CLM and RMS both designed the study, analyzed data, and wrote/edited the manuscript. FS and TS gathered/analyzed data and contributed toward writing/editing the manuscript.

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Data availability The datasets generated during and analyzed during the current study are available from the corresponding author on reasonable request.

Declarations

Conflict of interest RMS has received research support from Cerus, Grifols, and is part of a scientific advisory board for Octapharma. TS, FS, and CLM have no relevant disclosures.

Ethical approval This study was conducted under the oversight of the Emory University Institutional Review Board.

Informed consent The requirement for informed consent was waived by the Emory University Institutional Review Board.

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