



## THROMBOSIS AND HEMOSTASIS

# Rate of thrombosis in children and adolescents hospitalized with COVID-19 or MIS-C

Hilary Whitworth,<sup>1</sup> Sarah E. Sartain,<sup>2</sup> Riten Kumar,<sup>3</sup> Katherine Armstrong,<sup>4</sup> Lance Ballester,<sup>1</sup> Marisol Betensky,<sup>5,6</sup> Clay T. Cohen,<sup>2</sup> Rosa Diaz,<sup>2</sup> Caroline Diorio,<sup>1</sup> Neil A. Goldenberg,<sup>5-7</sup> Julie Jaffray,<sup>8</sup> Jacquelyn Keegan,<sup>8</sup> Kendra Malone,<sup>9</sup> Adrienne G. Randolph,<sup>10</sup> Stacey Rifkin-Zenenberg,<sup>4</sup> Wendy Seto Leung,<sup>8</sup> Anthony Sochet,<sup>11</sup> Lakshmi Srivaths,<sup>12</sup> Ayesha Zia,<sup>9</sup> and Leslie Raffini<sup>1</sup>

<sup>1</sup>Department of Pediatrics, Children's Hospital of Philadelphia, University of Pennsylvania, Philadelphia, PA; <sup>2</sup>Department of Pediatrics, Texas Children's Hospital, Baylor College of Medicine, Houston, TX; <sup>3</sup>Department of Pediatrics, Boston Children's Hospital, Harvard Medical School, Boston, MA; <sup>4</sup>Department of Pediatrics, Joseph M. Sanzari Children's Hospital, Hackensack University Medical Center, Hackensack, NJ; <sup>5</sup>Division of Hematology, Department of Pediatrics, Johns Hopkins University School of Medicine, Baltimore, MD; <sup>6</sup>Thrombosis Program and Institute for Clinical and Translational Research, Johns Hopkins All Children's Hospital, St Petersburg, FL; <sup>7</sup>Division of Hematology, Department of Medicine, Johns Hopkins University School of Medicine, Baltimore, MD; <sup>8</sup>Department of Pediatrics, Children's Hospital Los Angeles, University of Southern California Keck School of Medicine, Los Angeles, CA; <sup>9</sup>Department of Pediatrics, University of Texas Southwestern Medical Center, Dallas, TX; <sup>10</sup>Department of Anesthesiology, Critical Care and Pain Medicine, Boston Children's Hospital, Boston, MA; <sup>11</sup>Division of Pediatric Critical Care Medicine, Department of Anesthesia and Critical Care Medicine, Johns Hopkins University School of Medicine, Baltimore, MD; and <sup>12</sup>Department of Pediatrics, University of Texas Houston, Houston, TX

### KEY POINTS

- Cancer, central venous catheter, older age, and MIS-C are risk factors for thrombosis in children and adolescents with COVID-19 or MIS-C.
- Mortality was high (28%) in children and adolescents with MIS-C or COVID-19 who developed thrombosis.

**Coronavirus disease 2019 (COVID-19) caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) is associated with thrombotic complications in adults, but the incidence of COVID-19-related thrombosis in children and adolescents is unclear. Most children with acute COVID-19 have mild disease, but coagulopathy has been associated with multisystem inflammatory syndrome in children (MIS-C), a postinfectious complication. We conducted a multicenter retrospective cohort study to determine the incidence of thrombosis in children hospitalized with COVID-19 or MIS-C and evaluate associated risk factors. We classified patients into 1 of 3 groups for analysis: COVID-19, MIS-C, or asymptomatic SARS-CoV-2. Among a total of 853 admissions (COVID-19, n = 426; MIS-C, n = 138; and asymptomatic SARS-CoV-2, n = 289) in 814 patients, there were 20 patients with thrombotic events (TEs; including 1 stroke). Patients with MIS-C had the highest incidence (9 [6.5%] of 138) vs COVID-19 (9 [2.1%] of 426) or asymptomatic SARS-CoV-2 (2 [0.7%] of 289). In patients with COVID-19 or MIS-C, a majority of TEs (89%) occurred in patients age  $\geq 12$  years. Patients**

**age  $\geq 12$  years with MIS-C had the highest rate of thrombosis at 19% (9 of 48). Notably, 71% of TEs that were not present on admission occurred despite thromboprophylaxis. Multivariable analysis identified the following as significantly associated with thrombosis: age  $\geq 12$  years, cancer, presence of a central venous catheter, and MIS-C. In patients with COVID-19 or MIS-C, hospital mortality was 2.3% (13 of 564), but it was 28% (5 of 18) in patients with TEs. Our findings may help inform pediatric thromboprophylaxis strategies.**

## Introduction

The emergence of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) in late 2019 has led to the global pandemic of a highly transmissible and severe disease called coronavirus disease 2019 (COVID-19).<sup>1</sup> Although severe pneumonia is the cardinal presentation of COVID-19 in adults, a proclivity to cause thrombotic coagulopathy became apparent early in the pandemic.<sup>1-10</sup> Many patients had elevated levels of D-dimer and fibrinogen with mild thrombocytopenia and mild prolongation of prothrombin time (PT).<sup>1-3</sup> Studies in adults reported a high venous thromboembolism (VTE) rate, often occurring despite prophylactic anticoagulation.<sup>4-9</sup> A recent meta-analysis estimated an overall VTE incidence of 21% in adults hospitalized for

COVID-19, rising to 31% in those admitted to the intensive care unit (ICU).<sup>10</sup> The pooled odds of mortality were 74% higher among adult patients with thrombotic events (TEs), compared with those without TEs (23% vs 13%, respectively).<sup>10</sup>

Compared with adults, most children and adolescents with COVID-19 have minimal disease, and many infected with SARS-CoV-2 are asymptomatic.<sup>11</sup> Those age  $< 21$  years accounted for only 8% of reported cases and 0.08% of deaths in the United States.<sup>12</sup> Starting in April 2020, reports emerged of previously healthy children presenting with fever, cardiovascular shock and/or Kawasaki disease features, hyperinflammation, and multisystem involvement with a temporal association with SARS-CoV-2 exposure.<sup>13,14</sup> Many patients were SARS-CoV-2<sup>−</sup> on respiratory

testing but had positive antibody titers. Public health alerts from the Centers for Disease Control and Prevention (CDC) and the World Health Organization listed criteria for this new disease called multisystem inflammatory syndrome in children (MIS-C).<sup>15-17</sup> Coagulopathy was listed as a potential presenting feature of MIS-C.

As hospitalizations for COVID-19 increased, pediatric hematologists developed guidelines for thromboprophylaxis.<sup>18</sup> The Pediatric/Neonatal Scientific and Standardization Subcommittee of the International Society of Thrombosis and Haemostasis published consensus guidelines recommending pharmacologic prophylaxis in children with additional risk factors for VTE or those with elevated D-dimer.<sup>19</sup> These guidelines were based on extrapolation of adult data and expert opinion and not supported by data on the incidence of or risk factors for thrombosis in children or adolescents with COVID-19 or MIS-C. Therefore, the primary objective of this study was to determine the incidence of thrombotic complications in hospitalized children and adolescents with COVID-19 or MIS-C. We additionally sought to evaluate risk factors associated with TEs and describe the current practices for thromboprophylaxis in this cohort.

## Methods

### Study cohort

We conducted a retrospective cohort study of consecutive children age 0 to <21 years admitted from 1 March 2020 through 15 August 2020 with SARS-CoV-2<sup>+</sup> polymerase chain reaction (PCR) or diagnosis of MIS-C across 7 pediatric hospitals in 6 US states: Boston Children's Hospital (Massachusetts), Children's Hospital of Los Angeles (California), Children's Hospital of Philadelphia (Pennsylvania), Children's Medical Center of Dallas (Texas), Johns Hopkins All Children's Hospital (Florida), Joseph M. Sanzari Children's Hospital at Hackensack University Medical Center (New Jersey), and Texas Children's Hospital (Texas). This study was approved for a waiver of informed consent by each center's institutional review board.

Eligible patients were identified by  $\geq 1$  of the following: COVID-19 or MIS-C patient databases maintained at participating institutions, electronic medical record queries for patients with both an International Classification of Diseases (10th revision) COVID-19 diagnosis code and an inpatient admission code, and/or electronic medical record queries for patients with a SARS-CoV-2<sup>+</sup> test and coded for inpatient admission. Additionally, MIS-C patients were identified from electronic medical record queries for inpatients who had a consult order with the consult reason MIS-C and/or from electronic medical record queries for patients who had a COVID-19–related laboratory test and at least 1 of the following tests sent: ferritin, interleukin-2, or interleukin-6.

The following data were collected and entered into Research Electronic Data Capture software<sup>20,21</sup>: demographics, comorbidities, COVID-19 exposure history and presenting symptoms, hospital course (ICU admission; need for ventilator, vasopressors, or extracorporeal membrane oxygenation; presence of bacterial coinfection; presence of central venous catheter [CVC]; length of stay; and death), use of prophylactic anticoagulation (drug, dose, and duration), use of COVID-19– or MIS-C–directed therapies, hematologic and inflammatory laboratory parameters,

presence of so-called COVID toes,<sup>22</sup> presence of thrombosis up to 30 days postdischarge, and presence of bleeding during hospitalization.

### Study definitions

Admissions were categorized into 1 of 4 subgroups at the discretion of the investigator: COVID-19, a positive PCR and COVID-19 symptoms (fever, cough, diarrhea, loss of smell, headache, sore throat, or congestion); MIS-C, all 5 CDC criteria of fever, evidence of inflammation on laboratory testing,  $\geq 2$  organ systems involved, no other plausible diagnosis, and SARS-CoV-2<sup>+</sup> PCR or antibody test or known exposure<sup>23</sup>; MIS-C like, symptoms and laboratory findings highly suggestive of MIS-C in patients who were assessed and treated for MIS-C but did not fully meet the CDC criteria; and asymptomatic SARS-CoV-2, SARS-CoV-2<sup>+</sup> PCR test without symptoms of COVID-19 (listed above) and admitted with alternative diagnoses. Patients with asymptomatic SARS-CoV-2 were included primarily as a comparison group.

Clinically apparent thrombosis was defined as a radiologically confirmed arterial or venous thrombus. Superficial vein thrombosis was not included. In patients with thrombosis, data on clot location, symptoms, and anticoagulation treatment were collected. Bleeding was categorized as major or clinically relevant non-major based on the International Society of Thrombosis and Haemostasis consensus definition.<sup>24</sup> COVID-19–directed therapies included systemic steroids, convalescent plasma, remdesivir, tocilizumab, hydroxychloroquine, and anakinra. MIS-C–directed therapies included systemic steroids, IV immunoglobulin, anakinra, tocilizumab, and aspirin. Type, duration, and dosing of postdischarge thromboprophylaxis were collected. Laboratory parameters were collected as maximum and/or minimum values during hospitalization and included platelet count, hemoglobin, PT, lupus anticoagulant, D-dimer, and fibrinogen.

Obesity was defined using the CDC definition for body mass index  $\geq 95$ th percentile for age for children age  $> 24$  months.<sup>25</sup> If height and weight were not available, the patient was classified as obese if documented as such in the patient's medical record.

### Statistical analysis

Standard statistical methods were used to summarize the data: frequency and percentage for categorical variables, and median and interquartile range for continuous scaled variables. The incidence of TEs was calculated, along with the 95% confidence interval (CI) for each subgroup (COVID-19, MIS-C/MIS-C like, and asymptomatic SARS-CoV-2). Comparisons of demographic and clinical characteristics among patients with and without thrombosis were performed using nonparametric statistical tests. For univariable analyses, categorical variables were analyzed using count and percentage with Fisher's exact test, and continuous variables were analyzed using median, interquartile range, and Kruskal-Wallis rank test. Missing values were excluded from the analyses.

Multivariable analysis was performed using a binomial logit model with Wald test statistics. Given the low event rate (TEs), we only carried through a subset of the variables from the univariable analysis based on those that we believed to be independently associated. The primary analysis included patients

with COVID-19 or MIS-C/MIS-C like. Because of the low number of TEs, these groups were combined, and asymptomatic SARS-CoV-2 cases were excluded. A sensitivity analysis was performed excluding MIS-C-like patients.

All calculated *P* values were 2 sided, and an  $\alpha$  level of 0.05 was used for assessing significance; all analyses were conducted using SAS software (version 9.4).

## Results

Between 1 March and 15 August 2020, we identified 853 hospital admissions in 814 patients meeting eligibility criteria. Of these, 426 admissions (50%) were for COVID-19, 138 (16%) for MIS-C (*n* = 130) or MIS-C-like illness (*n* = 8), and 289 (34%) for asymptomatic SARS-CoV-2 infection. Of the 8 patients who did not meet full MIS-C criteria, 1 did not have documented fever, 5 did not have SARS-CoV-2<sup>+</sup> test or known exposure, and 2 had an alternative diagnosis. All had other features highly suggestive of MIS-C and were treated for MIS-C with steroids, IV immunoglobulin, aspirin, anakinra, and/or infliximab. Demographic and clinical characteristics of the cohort including therapies received are listed in Table 1 by clinical subgroup. For all subsequent analyses, MIS-C-like patients were combined with MIS-C patients. The MIS-C cohort had the highest proportion of patients without underlying comorbidities, with more evidence of coagulopathy (elevated D-dimer, fibrinogen, and PT and lower platelet count), and who required critical care (Table 1).

## TEs

There were 20 patients with  $\geq 1$  TE; 9 patients were admitted with COVID-19, 9 with MIS-C, and 2 with asymptomatic SARS-CoV-2. One patient admitted with COVID-19 developed a second TE during the admission (total, 21 TEs). Patients with MIS-C had the highest incidence at 6.5% (95% CI, 3% to 12%), followed by those with COVID-19 at 2.1% (95% CI, 1% to 4%) and those with asymptomatic SARS-CoV-2 at 0.7% (95% CI, 0.1% to 2.4%). A majority of TEs (89%) in COVID-19 or MIS-C patients occurred in those age  $\geq 12$  years. The incidence of TEs in patients age  $\geq 12$  years with COVID-19 or MIS-C was 6.8% (16 of 237), compared with 0.6% (2 of 327) in those age  $< 12$  years. In the subgroup of patients with MIS-C who were age  $\geq 12$  years, the incidence was 19% (9 of 48). Clinical details regarding patients with thrombosis are listed in Table 2. Of the 19 TEs in patients with COVID-19 or MIS-C, there were 11 deep vein thromboses, 3 pulmonary embolisms, 3 intracardiac thromboses, 1 acute ischemic stroke, and 1 cerebral sinovenous thrombosis. Four TEs were clinically apparent on admission to the hospital (Table 2). Thirteen (68%) were symptomatic, while the rest were asymptomatic, detected on imaging performed for another indication. Nine (47%) were catheter related. Death resulting from any cause occurred in 5 (28%) of the 18 patients with COVID-19 or MIS-C and thrombosis. A vast majority of COVID-19 and MIS-C patients with thrombosis (17 of 18) identified as being of either Hispanic ethnicity or African American race. One COVID-19 patient was diagnosed with COVID toes, but this was not included in overall TEs.

We evaluated several clinical, demographic, and laboratory risk factors for thrombosis in the combined cohort of COVID-19 and MIS-C, listed in Table 3. Older age, African American race and/or Hispanic ethnicity, MIS-C, admission to the ICU, need for a

ventilator, length of stay, presence of a CVC, cancer, D-dimer  $> 5 \times$  ULN, PT  $> 1.5 \times$  ULN, elevated fibrinogen, reduced platelet count, and death during admission were all statistically significant variables on univariable analysis. In a univariable analysis specifically comparing the rate of thrombosis in COVID-19 patients vs asymptomatic SARS-CoV-2 patients, COVID-19 was not significantly associated with TEs (*P* = .2).

## Data presented as *n* (%).

D-dimer  $> 5 \times$  ULN was significantly associated with TEs in the univariable analysis. However, there was a high proportion (36%) of missing D-dimer values, which was a problem for all the laboratory variables. Rather than imputing values, we ran the multivariable modeling in 2 ways: without laboratory values and including only admissions that had a D-dimer performed. In the model without laboratory values, the following variables were significantly associated with thrombosis: age  $\geq 12$  years, cancer, CVC, and MIS-C (Table 4). When D-dimer was included in the model, it was significant, but MIS-C was no longer statistically significant (Table 4).

In a sensitivity analysis that excluded the 8 MIS-C-like patients, the results of the multivariable analyses were similar to those of the primary analysis (data not shown).

## Thromboprophylaxis

Anticoagulant thromboprophylaxis was used during 30% (128 of 426) of COVID-19 and 58% (80 of 138) of MIS-C admissions, as well as during 61% of ICU admissions in those patients (Figure 1A). Nonetheless, 10 (71%) of the 14 TEs that were not identified on admission occurred in patients receiving thromboprophylaxis (regimens listed in Table 2). There were 220 regimens used in the 208 COVID-19 or MIS-C admissions where prophylactic anticoagulation was administered. The most common anticoagulant was enoxaparin (89%), followed by unfractionated heparin in 6.8% and other anticoagulants (4.5%). The variation with respect to drug and intensity is shown in Figure 1B. Once daily enoxaparin accounted for 20%, whereas a majority (62%) received twice daily prophylactic enoxaparin ( $\sim 0.5$  mg/kg every 12 hours). Full therapeutic intensity prophylaxis with enoxaparin ( $\sim 1$  mg/kg every 12 hours) or unfractionated heparin was used in 9.1%.

Notably, 45 (11%) of 417 patients with COVID-19 and 32 (25%) of 129 with MIS-C who did not develop thrombosis were discharged on prophylactic anticoagulation. A majority were discharged on enoxaparin (*n* = 56), followed by direct oral anticoagulants (*n* = 16), aspirin (*n* = 4), and warfarin (*n* = 1). The only patient known to develop thrombosis after discharge was a patient with cancer, found to have a catheter-associated intracardiac thrombosis within 30 days after an admission with asymptomatic SARS-CoV-2 when he was readmitted for COVID-19; this patient had not been discharged on anticoagulation.

## Bleeding

There were 9 admissions with major bleeding and 8 with clinically relevant nonmajor bleeding among the COVID-19 and MIS-C subgroups (Table 1). The COVID-19 cohort had 7 admissions (1.6%) with major bleeding events, 2 of which occurred while on anticoagulation, and there were 2 (1.4%) in the MIS-C cohort with major bleeding (both events occurred on anticoagulation). Of the 4 COVID-19 or MIS-C patients with major bleeding on

**Table 1. Clinical characteristics of all 3 SARS-CoV-2 cohorts**

	Asymptomatic SARS-CoV-2 (n = 289)	COVID-19 (n = 426)	MIS-C/ MIS-C like (n = 138)
<b>Age, y</b>			
Median	10	10	10
IQR	3-15	1-16	6-14
Male sex	144 (50)	226 (53)	75 (54)
<b>Ethnicity</b>			
Hispanic	125 (43)	228 (54)	57 (41)
Non-Hispanic	139 (48)	167 (39)	69 (50)
Unknown/other	25 (9)	31 (7)	12 (9)
<b>Race</b>			
White	151 (52)	234 (55)	77 (56)
African American	74 (26)	89 (21)	39 (28)
Asian	2 (1)	6 (1)	4 (3)
Other/unknown	62 (21)	97 (23)	18 (13)
African American race and/or Hispanic ethnicity	198 (69)	315 (74)	96 (70)
<b>Length of stay, d</b>			
Median	3	3	6
IQR	1-7	2-7	4-10
<b>Comorbidity</b>			
Respiratory	22 (8)	90 (21)	16 (12)
Cardiac	13 (5)	38 (9)	5 (4)
Cancer	25 (9)	39 (9)	5 (4)
Obesity	61 (21)	114 (27)	45 (33)
Diabetes	17 (6)	15 (4)	0
Immunodeficiency	10 (3)	18 (4)	1 (1)
Sickle cell disease	8 (3)	26 (6)	1 (1)
Neurologic	38 (13)	49 (12)	4 (3)
Gastrointestinal	36 (12)	38 (9)	1 (1)
None	77 (27)	114 (27)	70 (51)
<b>ICU admission</b>			
Ventilator	46 (16)	136 (32)	86 (62)
ECMO	11 (4)	46 (11)	22 (16)
	1 (0.4)	4 (1)	3 (2)
CVC	47 (16)	74 (17)	53 (38)
COVID-19-directed treatment	0	105 (25)	39 (28)
MIS-C-directed treatment	1 (0.3)	6 (1)	130 (94)
<b>Maximum D-dimer</b>			
Not done	214 (74)	197 (46)	5 (4)
Normal range	14 (5)	36 (8)	1 (1)
1-5× ULN	34 (12)	131 (31)	32 (23)
>5× ULN	27 (9)	62 (15)	100 (72)
<b>Minimum platelet count, ×10<sup>3</sup>/μL</b>			
Median	259	208	148
IQR	199-320	140-273	103-204

**Table 1. (continued)**

	Asymptomatic SARS-CoV-2 (n = 289)	COVID-19 (n = 426)	MIS-C/ MIS-C like (n = 138)
<b>Maximum fibrinogen, mg/dL</b>			
Median	421	463	551
IQR	282-583	325-583	462-657
<b>Maximum PT</b>			
Not done	183 (63)	204 (48)	7 (5)
Normal range	55 (19)	121 (28)	38 (27)
1-1.5× ULN	48 (17)	90 (21)	86 (62)
>1.5× ULN	3 (1)	11 (3)	7 (5)
Thromboprophylaxis	31 (11)	128 (30)	80 (58)
<b>Admission with TEs</b>			
DVT	2 (0.7)	9 (2)	9 (7)
PE	2	4	7
Stroke	0	3	0
Intracardiac	0	0	1
Cerebral sinovenous thrombosis	0	2	1
	0	1	0
Major bleeding	4 (1)	7 (2)	2 (1)
Death during hospitalization (or <30 d)	2 (0.7)	11 (2.6)	2 (1.4)

Data presented as n (%).

CVC, central venous catheter; DVT, deep vein thrombosis; ECMO, extracorporeal membrane oxygenation; IQR, interquartile range; PE, pulmonary embolism; ULN, upper limit of normal.

anticoagulation, 1 was receiving prophylactic daily enoxaparin, 1 was receiving therapeutic-dose unfractionated heparin for extracorporeal membrane oxygenation and treatment of TEs, and 2 were receiving therapeutic-dose enoxaparin for treatment of TEs.

## Mortality

The all-cause mortality rate in our cohort of COVID-19 and MIS-C patients was 2.3% (13 of 564). A majority of deaths (n = 11) occurred in patients hospitalized with COVID-19. All patients who died had an underlying comorbidity; the most common was cancer (n = 7). Respiratory failure was the most common reason for death (n = 7). The cause of death was attributed, at least in part, to malignancy in 3 patients. One patient had a hemothorax while on therapeutic anticoagulation that was thought to contribute to death. Other reported contributors to death included renal failure (n = 1), thrombotic microangiopathy (n = 1), complex congenital heart disease (n = 1), bowel obstruction with abdominal compartment syndrome (n = 1), and septic shock (n = 1). Five of the 18 patients (28%) with COVID-19 or MIS-C and TEs died.

## Discussion

This is the first study to evaluate the incidence of thrombosis in a large, multicenter cohort of children hospitalized with



**Table 2. Clinical characteristics of patients who developed thrombosis**

Patient	Age, y*	Sex	Race/ethnicity	Thrombosis location	Catheter related	Underlying medical conditions	Thromboprophylaxis	ICU	Other†
<b>COVID-19</b>	1	F	White/Hispanic	PE	No	Cancer (relapsed sarcoma), CVC, obesity	Enoxaparin 30 mg every 12 h	Yes	Critically ill; died as a result of cardiac arrest, COVID-19 pneumonia, hemothorax
	2	M	African American	Upper-extremity DVT	Yes	Congenital heart disease, cerebral palsy, CVC, bacterial tracheitis	Enoxaparin every 12 h, goal anti-Xa 0.2-0.5 IU/mL	Yes	Critically ill; died as a result of cardiac arrest
	3	M	White/Hispanic	PE	No	Cancer (AML), CVC	No; thrombosis present on admission	Yes	Critically ill; died as a result of intracranial hemorrhage, COVID-19, multisystem organ failure
	4	M	White/Hispanic	Lower-extremity DVT	No	Obesity	No	Yes	Died as a result of AML
	5	F	White/Hispanic	Intracardiac thrombosis	Yes	Cancer (AML), CVC	No	Yes	Critically ill
	6	F	African American	PE	No	Obesity	Enoxaparin 0.5 mg/kg every 12 h	Yes	Major bleeding event on anticoagulation
	7	M	White/Hispanic	Cerebral sinovenous thrombosis	No	Cancer	No	No	
	8	F	African American	Upper-extremity DVT ×2	Yes	Cancer, obesity, CVC	No; thrombosis present on admission, second thrombosis occurred while on enoxaparin 1 mg/kg every 12 h	No	
	9	M	Other/Hispanic	Intracardiac thrombosis	Yes	Cancer, CVC	No; thrombosis present on admission	No	Readmitted after asymptomatic SARS-CoV-2 admission with symptomatic COVID-19, thrombosis on chest CT
<b>MIS-C</b>	10	M	African American	Acute ischemic stroke	No	Neurologic disorder, CVC	No; stroke on admission	Yes	Critically ill; admitted with acute MCA stroke and MIS-C 1 mo after acute COVID-19 admission
	11	F	African American	Extremity DVT	No	Respiratory, CVC	No	Yes	Critically ill; admitted in shock and with decreased cardiac function
	12‡	F	Unknown	Lower-extremity DVT	No	None, CVC	Enoxaparin every 12 h, goal anti-Xa 0.3-0.5 IU/mL	Yes	Critically ill, on ECMO
	13	M	African American	Lower-extremity DVT	No	None, CVC	UFH, goal anti-Xa 0.1-0.3 IU/mL	Yes	Critically ill
	14	M	Unknown/Hispanic	Intracardiac thrombosis	Yes	Obesity, CVC	UFH, goal anti-Xa 0.3-0.7 IU/mL	Yes	Critically ill; thrombosis found post-ECMO
	15	F	African American	Upper-extremity DVT	Yes	Obesity, CVC	Heparin 5000 units every 12 h then enoxaparin, goal anti-Xa 0.5-1.0 IU/mL	Yes	Critically ill
	16‡	F	White/Hispanic	Upper-extremity DVT	Yes	Cancer, obesity, respiratory, CVC	Enoxaparin daily	No	Died as a result of multiorgan failure
	17‡	M	African American	Upper-extremity DVT	No	Cancer, obesity, CVC	Enoxaparin, goal anti-Xa 0.3-0.5 IU/mL	No	Symptomatic COVID-19 then readmitted with MIS-C, relapsed ALL, bacteremia
	18	F	African American	Upper-extremity DVT	Yes	Obesity, CVC	Enoxaparin, goal anti-Xa 0.2-0.5 IU/mL	No	Urinary tract infection
<b>Asymptomatic SARS-CoV-2</b>	19	F	White/Hispanic	Lower-extremity DVT	No	Cancer (ALL in induction)	Enoxaparin 40 mg daily	Yes	Died as a result of complications of ALL; bacteremia
	20	F	African American	Lower-extremity DVT	Yes	Acute liver failure, CVC	No	Yes	

ALL, acute lymphoblastic leukemia; AML, acute myeloid leukemia; CT, computed tomography; CVC, central venous catheter; DVT, deep vein thrombosis; ECMO, extracorporeal membrane oxygenation; MCA, middle cerebral artery; PE, pulmonary embolism; UFH, unfractionated heparin.

\*Age range presented so patient identification less likely.

†Critically ill indicates patient required mechanical ventilation, vasopressors, and/or ECMO.

‡MIS-C like.

**Table 3. Comparison of COVID-19 or MIS-C/MIS-C-like patients with thrombosis with those without**

	Thrombosis (n = 18)	No thrombosis (n = 546)	P
<b>Age, y</b>			.0001
Median	16	10	
IQR	14-17	2-15	
<b>Age group, y</b>			.0002
0-6	1 (6)	209 (38)	
6-12	1 (6)	116 (21)	
≥12	16 (89)	221 (40)	
Male sex	9 (50)	292 (53)	.81
Hispanic ethnicity	8 (44)	277 (51)	.64
<b>Race</b>			
White	6 (33)	305 (56)	.09
African American	9 (50)	119 (22)	.009
Asian	0	10 (2)	1
Other/unknown	3 (17)	112 (21)	1
African American race and/or Hispanic ethnicity	17 (94)	394 (72)	.03
<b>SARS-CoV-2 disease category</b>			.02
COVID-19	9 (50)	417 (76)	
MIS-C	9 (50)	129 (24)	
<b>Length of stay, d</b>			<.0001
Median	18	4	
IQR	10-41	2-8	
<b>Comorbidity</b>			
Respiratory	3 (17)	103 (19)	1
Cardiac	1 (6)	42 (8)	1
Cancer	8 (44)	36 (7)	<.0001
Obesity	9 (50)	150 (27)	.06
Neurologic	2 (11)	51 (9)	.68
Sickle cell disease	0	27	1
None	2 (11)	182 (33)	.07
<b>ICU admission</b>			.006
Ventilator	9 (50)	59 (11)	.004
ECMO	2 (11)	5 (1)	.06
CVC	15 (83)	112 (21)	<.0001
<b>Maximum D-dimer</b>			.002
Not done	2 (11)	200 (37)	
Normal range	0	37 (7)	
1-5× ULN	2 (11)	161 (29)	
>5× ULN	14 (78)	148 (27)	
<b>Maximum fibrinogen, mg/dL</b>			.034
Median	632	499	
IQR	449-788	386-602	
<b>Minimum platelet count, ×10<sup>3</sup>/μL</b>			.0001
Median	71	189	
IQR	54-152	131-266	

**Table 3. (continued)**

	Thrombosis (n = 18)	No thrombosis (n = 546)	P
<b>PT</b>			.03
Not done	1 (6)	210 (38)	
Normal range	3 (17)	156 (29)	
1-1.5× ULN	12 (67)	164 (30)	
>1.5× ULN	2 (11)	16 (3)	
Thromboprophylaxis	10 (56)	198 (36)	.134
Bacterial coinfection	5 (28)	67 (12)	.07
Death during hospitalization (or <30 d)	5 (28)	8 (1)	<.0001

Data presented as n (%).

CVC, central venous catheter; ECMO, extracorporeal membrane oxygenation; IQR, interquartile range; PT, prothrombin time.

COVID-19–related complications.<sup>16,26-28</sup> The incidence of TEs in hospitalized children was 2.1% in those with COVID-19 and 6.5% in those with MIS-C, compared with 0.7% in those with asymptomatic SARS-CoV-2 infection. We identified several important laboratory (elevated D-dimer) and clinical variables (age ≥12 years, cancer, MIS-C, and CVC) that were associated with a much higher risk of TEs. Importantly, more than two-thirds of TEs that occurred in hospitalized children with COVID-19 or MIS-C occurred in patients receiving thromboprophylaxis. These findings have important implications for clinicians and provide new evidence regarding the risk of TEs in children hospitalized with COVID-19 or MIS-C. Mortality was 28% in pediatric patients with TEs and COVID-19 or MIS-C, although most patients who died had comorbid conditions that were risk factors for TEs and contributed to mortality.<sup>10</sup>

To put our findings into perspective, the rate of VTE in children admitted to US tertiary care hospitals in 2007 was estimated at 58 (0.58%) per 10 000 admissions using the Pediatric Health Information System database, with those age 12 to 18 years having the highest rate (94 [0.94%] per 10 000).<sup>29</sup> The rate observed in our study was much higher, particularly in patients age ≥12 years (6.8%) with COVID-19 or MIS-C. However, this comparison is limited for several reasons. Patients hospitalized with COVID-19 are known to have a high prevalence of underlying medical conditions (eg, cancer, diabetes, and obesity), which may have increased the rate. In addition, the age distribution of patients in this study was skewed toward older patients compared with the Pediatric Health Information System study, which would also result in a higher rate. Nonetheless, the rates of TEs that we observed, particularly in the MIS-C population and those age ≥12 years, suggest that COVID-19 and MIS-C are unique risk factors for thrombosis in hospitalized children.

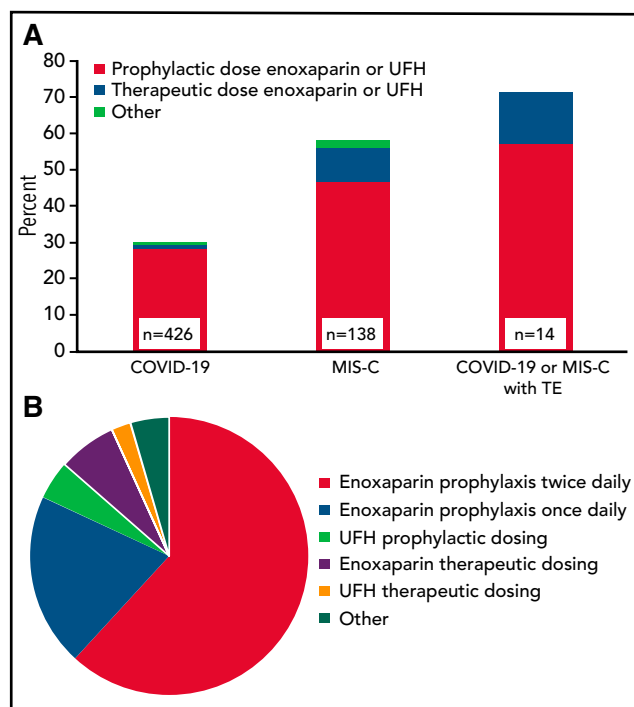
Consistent with this conclusion is the finding that a D-dimer >5× ULN was significantly associated with TEs in our study. This is the first pediatric COVID-19 study to demonstrate the association between elevated D-dimer and TEs, and this finding is consistent with multiple studies of adults with COVID-19.<sup>8,30,31</sup> In addition, elevated fibrinogen, prolonged PT, and reduced

**Table 4. Multivariable model for factors associated with thrombosis in COVID-19 and MIS-C/MIS-C-like patients**

Variable	Without D-dimer		With D-dimer*	
	OR (95% CI)	P	OR (95% CI)	P
Cancer	6.34 (1.56-25.73)	.01	13.74 (2.36-79.95)	.004
Obesity	0.89 (0.24-3.25)	.856	1.16 (0.21-6.35)	.863
No comorbidity	0.31 (0.04-2.17)	.236	0.43 (0.05-3.79)	.447
Bacterial coinfection	1.63 (0.41-6.48)	.484	1.31 (0.27-6.43)	.741
CVC	7.22 (1.71-30.45)	.007	25.71 (2.65-249.01)	.005
ICU admission	2.17 (0.56-8.40)	.263	0.35 (0.04-3.04)	.339
MIS-C	6.44 (1.65-25.24)	.008	3.18 (0.57-17.55)	.185
COVID-19	Reference		Reference	
Age <6 y	1.30 (0.06-27.71)	.871	1.19 (0.05-30.28)	.916
Age 6-12 y	Reference		Reference	
Age ≥12 y	16.84 (1.93-147.1)	.011	20.05 (2.18-184.29)	.008
African American race and/or Hispanic ethnicity	7.14 (0.83-61.36)	.073	9.74 (0.91-104.45)	.06
D-dimer >5× ULN	Not included		21.16 (2-223.94)	.011

CVC, central venous catheter; OR, odds ratio; ULN, upper limit of normal

\*Missing values excluded.



**Figure 1. Prophylactic anticoagulation regimens in COVID-19 and MIS-C patients.** (A) Proportion of admissions receiving prophylactic anticoagulation in clinical subgroups by dose intensity (n = 208 admissions). When >1 regimen was used per admission, the regimen with either the longest duration or highest dose intensity was included. For patients with TEs, only those without TEs on admission were included (n = 14). (B) Prophylactic anticoagulation regimens in patients with COVID-19 or MIS-C (n = 220 regimens; other: rivaroxaban, apixaban, bivalirudin, warfarin, or aspirin). TE, thrombotic event; UFH, unfractionated heparin.

platelet count were also associated with TEs in our study on univariable analysis, similar to reports of adults with COVID-19.<sup>1-3</sup> A limitation of our analysis is that there was a high proportion of missing laboratory values, and we limited the collection of laboratory parameters to the maximum or minimum values during hospitalization, regardless of timing (ie, early in hospitalization and/or before TEs). Therefore, it is possible that elevations in D-dimer levels may have occurred after a TE in some patients, reducing the predictive value of this variable. Future studies will be necessary to better evaluate the predictive value of elevated D-dimer in this setting.

In a multivariable regression model that excluded laboratory values, MIS-C, age ≥12 years, presence of a CVC, and cancer were significantly associated with TEs. With the exception of MIS-C, these mirror previously established risk factors for TEs in hospitalized children.<sup>32,33</sup> When D-dimer was included in the multivariable model, it was significant; however, MIS-C was no longer significant. We hypothesize that this is due to the collinearity of these variables, as well as to the proportion of missing laboratory values. Interestingly, cancer was identified as an independent risk factor for TEs in multivariable models with and without D-dimer. Although COVID-19 was not found to be a risk factor for TEs, 30% of patients with COVID-19 received thromboprophylaxis, which may have reduced the rate of thrombosis in this group. In addition, our ability to evaluate COVID-19 as a risk factor may have been limited by the small sample size. Because of the small sample size, missing laboratory values, and lack of a validation cohort, we did not develop a clinical prediction model. Future studies, focused on TE risk prediction in children with SARS-CoV-2, should address these points and also include other potentially important variables, such as blood type and pubertal status (rather than age).

We decided to include asymptomatic patients with SARS-CoV-2 for several reasons. First, we wanted to evaluate the risk of thrombosis in these children, because hematologists are frequently consulted to address the role of thromboprophylaxis in this setting. Second, if patients were truly asymptomatic and had no effects from viremia, this group could potentially serve as a control group. The overall incidence of TEs in the asymptomatic SARS-CoV-2 group was low (0.7%), and the 2 patients with TEs in this subgroup had several prothrombotic risk factors (Table 2). Therefore, we believe that asymptomatic SARS-CoV-2 infection may not significantly increase the risk of thrombosis.

In our study, all 9 of the TEs in the MIS-C cohort occurred in patients age  $\geq 12$  years, for a rate of 19% (9 of 48) in this subgroup. This is higher than that in an earlier report from a multicenter registry of 186 MIS-C patients, where 3 (6.7%) of 45 patients age  $> 12$  years had TEs.<sup>16</sup> This difference may be due to variation in thromboprophylaxis, severity of disease, inclusion of different centers, or because our study was designed to focus on TEs. We did not identify any TEs in the 90 MIS-C admissions that occurred in children age  $< 12$  years, confirming the low incidence of TEs in younger children with MIS-C observed in the prior study.

Similar to other studies, we observed a high prevalence of Hispanic (51%) and African American children (23%) hospitalized with COVID-19 or MIS-C.<sup>34</sup> Of the 18 patients with TEs and COVID-19 or MIS-C, 17 identified as being of either Hispanic ethnicity or African American race. These racial disparities associated with worse outcomes in both children and adults are likely linked to similar socioeconomic disadvantages that have been suggested by others.<sup>12,34,35</sup>

A majority of TEs in our study developed while patients were receiving thromboprophylaxis, which is similar to adult reports.<sup>7,8</sup> This has raised questions regarding the optimal intensity of prophylactic anticoagulation for patients with SARS-CoV-2. We are unable to answer this question because thromboprophylactic practices varied across institutions and changed during the study period. However, it is notable that 4 of 9 of the major bleeding events in patients with COVID-19 and MIS-C occurred on anticoagulation. This is limited by the fact that we could not assess all clinical risk factors for bleeding in these patients; however, anticoagulation likely contributed to these bleeding events. Optimal thromboprophylactic dose intensity may be more carefully addressed by an ongoing clinical trial to evaluate the safety of enoxaparin in children with SARS-CoV-2-associated illness, administered twice daily to achieve a 4-hour postdose anti-Xa level of 0.20 to 0.49 IU/mL.<sup>36</sup>

There are several limitations to this retrospective study. First, our findings must be contextualized, recognizing that a significant proportion of patients received inpatient and postdischarge thromboprophylaxis. Given the lack of a comparator group, it is plausible that the rate of thrombosis in this cohort would have been higher in the absence of thromboprophylaxis. Second, participating hospitals were large pediatric referral centers, and patients may have had more medically complex cases or been transferred from outside hospitals for a higher level of care. Thus, the risk of thrombosis may actually be an overestimate, and results from our study may not be generalizable to smaller pediatric centers or community hospitals. In contrast, some

patients may have died as a result of unrecognized TEs or PE; this number is likely low, but would have resulted in an underestimate of the risk. Third, there was some variation over time regarding the practice of testing all admitted patients for SARS-CoV-2 (particularly earlier in the pandemic), so patients with mild disease may have been missed. Last, it is important to note that a proportion of patients in this study (estimated at  $< 10\%$ ) have been reported in prior publications or included in other registries.<sup>16,37-39</sup> To our knowledge, none of the TEs have been previously reported.

For the first time, we provide data regarding the rate of thrombotic complications in patients age  $< 21$  years hospitalized with SARS-CoV-2-associated illness. Our findings, including the low rate of TEs in children age  $< 12$  years with COVID-19 or MIS-C and the low rate of postdischarge TEs, along with factors associated with an increased risk (age  $\geq 12$  years, MIS-C, CVC, and cancer), may help inform thromboprophylactic strategies at pediatric centers. Future studies focused on the intensity of anticoagulation as well as on novel targets are well under way in adults with COVID-19 and will likely be relevant to high-risk adolescent patients with COVID-19 or MIS-C, in whom the rates of thrombosis and mortality were highest.

## Acknowledgments

This study was supported by National Institutes of Health, National Heart, Lung, and Blood Institute grant T32 HL007971 (H.W.) and a scholarship from the Institute for Translational Medicine and Therapeutics at the University of Pennsylvania (C.D.).

## Authorship

Contribution: H.W., S.E.S., R.K., K.A., L.B., M.B., C.T.C., R.D., C.D., N.A.G., J.J., A.G.R., S.R.-Z., A.S., L.S., A.Z., and L.R. contributed to study design; H.W., S.E.S., R.K., K.A., M.B., C.T.C., R.D., C.D., J.J., J.K., K.M., S.R.-Z., W.S.L., A.S., L.S., A.Z., and L.R. contributed to data collection; H.W., L.B., and L.R. analyzed data; and H.W., S.E.S., R.K., K.A., L.B., M.B., C.T.C., R.D., C.D., N.A.G., J.J., A.G.R., S.R.-Z., A.S., L.S., A.Z., and L.R. wrote and edited the paper.

Conflict-of-interest disclosure: S.E.S. serves on the advisory board for Alexion Pharmaceuticals. R.K. has attended advisory board meetings for Bayer, Genentech, and Kedrion. The remaining authors declare no competing financial interests.

ORCID profiles: H.W., 0000-0002-5323-7023; C.T.C., 0000-0001-6819-4049; C.D., 0000-0002-8005-3836; J.J., 0000-0002-1175-7266; A.G.R., 0000-0002-3084-3071; A.Z., 0000-0003-3283-0415; L.R., 0000-0001-8152-1040.

Correspondence: Leslie Raffini, 11022 Colkett Translational Research Building, 3501 Civic Center Blvd, Philadelphia, PA 19104; e-mail: raffini@chop.edu.

## Footnotes

Submitted 5 January 2021; accepted 30 January 2021; prepublished online on *Blood* First Edition 25 April 2021. DOI 10.1182/blood.2020010218.

For original data, please contact raffini@chop.edu.

There is a *Blood* Commentary on this article in this issue.

The publication costs of this article were defrayed in part by page charge payment. Therefore, and solely to indicate this fact, this article is hereby marked "advertisement" in accordance with 18 USC section 1734.



## REFERENCES

- Zhou F, Yu T, Du R, et al. Clinical course and risk factors for mortality of adult inpatients with COVID-19 in Wuhan, China: a retrospective cohort study. *Lancet*. 2020; 395(10229):1054-1062.
- Fogarty H, Townsend L, Ni Cheallagh C, et al. COVID19 coagulopathy in Caucasian patients. *Br J Haematol*. 2020;189(6):1044-1049.
- Wang D, Hu B, Hu C, et al. Clinical characteristics of 138 hospitalized patients with 2019 novel coronavirus-infected pneumonia in Wuhan, China. *JAMA*. 2020; 323(11):1061-1069.
- Al-Samkari H, Karp Leaf RS, Dzik WH, et al. COVID-19 and coagulation: bleeding and thrombotic manifestations of SARS-CoV-2 infection. *Blood*. 2020;136(4):489-500.
- Cui S, Chen S, Li X, Liu S, Wang F. Prevalence of venous thromboembolism in patients with severe novel coronavirus pneumonia. *J Thromb Haemost*. 2020;18(6):1421-1424.
- Helms J, Tacquard C, Severac F, et al; CRICS TRIGGERSEP Group (Clinical Research in Intensive Care and Sepsis Trial Group for Global Evaluation and Research in Sepsis). High risk of thrombosis in patients with severe SARS-CoV-2 infection: a multicenter prospective cohort study. *Intensive Care Med*. 2020;46(6):1089-1098.
- Litjens JF, Leclerc M, Chochois C, et al. High incidence of venous thromboembolic events in anticoagulated severe COVID-19 patients. *J Thromb Haemost*. 2020;18(7):1743-1746.
- Middelorp S, Coppens M, van Haaps TF, et al. Incidence of venous thromboembolism in hospitalized patients with COVID-19. *J Thromb Haemost*. 2020;18(8):1995-2002.
- Poissy J, Goutay J, Caplan M, et al; Lille ICU Haemostasis COVID-19 Group. Pulmonary embolism in patients with COVID-19: awareness of an increased prevalence. *Circulation*. 2020;142(2):184-186.
- Malas MB, Naazie IN, Elsayed N, Mathlouthi A, Marmor R, Clary B. Thromboembolism risk of COVID-19 is high and associated with a higher risk of mortality: a systematic review and meta-analysis. *EClinicalMedicine*. 2020; 29:100639.
- Lu X, Zhang L, Du H, et al; Chinese Pediatric Novel Coronavirus Study Team. SARS-CoV-2 infection in children. *N Engl J Med*. 2020; 382(17):1663-1665.
- Bixler D, Miller AD, Mattison CP, et al; Pediatric Mortality Investigation Team. SARS-CoV-2-associated deaths among persons aged <21 years - United States, February 12-July 31, 2020. *MMWR Morb Mortal Wkly Rep*. 2020;69(37):1324-1329.
- Riphagen S, Gomez X, Gonzalez-Martinez C, Wilkinson N, Theocharis P. Hyperinflammatory shock in children during COVID-19 pandemic. *Lancet*. 2020;395(10237):1607-1608.
- Verdoni L, Mazza A, Gervasoni A, et al. An outbreak of severe Kawasaki-like disease at the Italian epicentre of the SARS-CoV-2 epidemic: an observational cohort study. *Lancet*. 2020;395(10239):1771-1778.
- Dufort EM, Koumans EH, Chow EJ, et al; New York State and Centers for Disease Control and Prevention Multisystem Inflammatory Syndrome in Children Investigation Team. Multisystem inflammatory syndrome in children in New York state. *N Engl J Med*. 2020;383(4): 347-358.
- Feldstein LR, Rose EB, Horwitz SM, et al; CDC COVID-19 Response Team. Multisystem inflammatory syndrome in U.S. children and adolescents. *N Engl J Med*. 2020;383(4):334-346.
- Ramcharan T, Nolan O, Lai CY, et al. Paediatric inflammatory multisystem syndrome: temporally associated with SARS-CoV-2 (PIMS-TS): cardiac features, management and short-term outcomes at a UK tertiary paediatric hospital. *Pediatr Cardiol*. 2020;41(7):1391-1401.
- Loi M, Branchford B, Kim J, Self C, Nuss R. COVID-19 anticoagulation recommendations in children. *Pediatr Blood Cancer*. 2020;67(9): e28485.
- Goldenberg NA, Sochet A, Albisetti M, et al; Pediatric/Neonatal Hemostasis and Thrombosis Subcommittee of the ISTH SSC. Consensus-based clinical recommendations and research priorities for anticoagulant thromboprophylaxis in children hospitalized for COVID-19-related illness. *J Thromb Haemost*. 2020;18(11):3099-3105.
- Harris PA, Taylor R, Thielke R, Payne J, Gonzalez N, Conde JG. Research electronic data capture (REDCap)—a metadata-driven methodology and workflow process for providing translational research informatics support. *J Biomed Inform*. 2009;42(2): 377-381.
- Harris PA, Taylor R, Minor BL, et al; REDCap Consortium. The REDCap Consortium: building an international community of software platform partners. *J Biomed Inform*. 2019;95:103208.
- Cordoro KM, Reynolds SD, Wattier R, McCalmont TH. Clustered cases of acral pernio: clinical features, histopathology, and relationship to COVID-19. *Pediatr Dermatol*. 2020;37(3):419-423.
- Centers for Disease Control and Prevention. Multisystem inflammatory syndrome in children (MIS-C) associated with coronavirus disease 2019 (COVID-19). <https://emergency.cdc.gov/han/2020/han00432.asp>. Accessed 30 July 2020.
- Mitchell LG, Goldenberg NA, Male C, Kenet G, Monagle P, Nowak-Göttl U; Perinatal and Paediatric Haemostasis Subcommittee of the SSC of the ISTH. Definition of clinical efficacy and safety outcomes for clinical trials in deep venous thrombosis and pulmonary embolism in children. *J Thromb Haemost*. 2011;9(9):1856-1858.
- Abarca-Gómez L, Abdeen ZA, Hamid ZA, et al; NCD Risk Factor Collaboration (NCD-RisC). Worldwide trends in body-mass index, underweight, overweight, and obesity from 1975 to 2016: a pooled analysis of 2416 population-based measurement studies in 128.9 million children, adolescents, and adults. *Lancet*. 2017;390(10113):2627-2642.
- Ippolito Bastidas H, Márquez-Pérez T, García-Salido A, et al. Cerebral venous sinus thrombosis in a pediatric patient with COVID-19. *Neurol Clin Pract*. 2020;11(2): e208-e210.
- Essajee F, Solomons R, Goussard P, Van Toorn R. Child with tuberculous meningitis and COVID-19 coinfection complicated by extensive cerebral sinus venous thrombosis. *BMJ Case Rep*. 2020;13(9):e238597.
- Del Borrello G, Giraudo I, Bondone C, et al. SARS-CoV-2 associated coagulopathy and thromboembolism prophylaxis in children: a single centre observational study. *J Thromb Haemost*. 2021;19(2):522-530.
- Raffini L, Huang Y-S, Witmer C, Feudtner C. Dramatic increase in venous thromboembolism in children's hospitals in the United States from 2001 to 2007. *Pediatrics*. 2009; 124(4):1001-1008.
- Miesbach W, Makris M. COVID-19: coagulopathy, risk of thrombosis, and the rationale for anticoagulation. *Clin Appl Thromb Hemost*. 2020;26:1076029620938149.
- Zhang L, Feng X, Zhang D, et al. Deep vein thrombosis in hospitalized patients with coronavirus disease 2019 (COVID-19) in Wuhan, China: prevalence, risk factors, and outcome [published correction appears in *Circulation*. 2020;142(2):e33]. *Circulation*. 2020;142(2): 114-128.
- Mahajerin A, Branchford BR, Amankwah EK, et al. Hospital-associated venous thromboembolism in pediatrics: a systematic review and meta-analysis of risk factors and risk-assessment models. *Haematologica*. 2015;100(8):1045-1050.
- Witmer CM, Takemoto CM. Pediatric hospital acquired venous thromboembolism. *Front Pediatr*. 2017;5:198.
- Goyal MK, Simpson JN, Boyle MD, et al. Racial and/or ethnic and socioeconomic disparities of SARS-CoV-2 infection among children. *Pediatrics*. 2020;146(4): e202009951.
- Alcendor DJ. Racial disparities-associated COVID-19 mortality among minority populations in the US. *J Clin Med*. 2020; 9(8):2442.
- COVID-19 anticoagulation in children - thromboprophylaxis. <https://clinicaltrials.gov/ct2/show/NCT04354155>.
- Diorio C, Anderson EM, McNerney KO, et al. Convalescent plasma for pediatric patients with SARS-CoV-2-associated acute respiratory distress syndrome. *Pediatr Blood Cancer*. 2020;67(11):e28693.
- Chiotos K, Bassiri H, Behrens EM, et al. Multisystem inflammatory syndrome in children during the COVID-19 pandemic: a case series. *J Pediatric Infect Dis Soc*. 2020; 9(3):393-398.
- Diorio C, McNerney KO, Lambert M, et al. Evidence of thrombotic microangiopathy in children with SARS-CoV-2 across the spectrum of clinical presentations. *Blood Adv*. 2020;4(23):6051-6063.