



Since January 2020 Elsevier has created a COVID-19 resource centre with free information in English and Mandarin on the novel coronavirus COVID-19. The COVID-19 resource centre is hosted on Elsevier Connect, the company's public news and information website.

Elsevier hereby grants permission to make all its COVID-19-related research that is available on the COVID-19 resource centre - including this research content - immediately available in PubMed Central and other publicly funded repositories, such as the WHO COVID database with rights for unrestricted research re-use and analyses in any form or by any means with acknowledgement of the original source. These permissions are granted for free by Elsevier for as long as the COVID-19 resource centre remains active.



Acute limb ischemia among patients with COVID-19 infection

George Galyfos, MD, PhD,^a Argiri Sianou, MD,^b Maximos Frountzas, MD,^a Kotsarinis Vasilios, MD,^a Dimitrios Vouros, MD,^a Charis Theodoropoulos, MD,^a Victoria Michalopoulou, MD,^a Frangiska Sigala, MD, PhD,^a and Konstantinos Filis, MD, PhD,^a Athens, Greece

ABSTRACT

Objective/Background: Severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) infection has been associated with thrombotic complications such as deep vein thrombosis or stroke. Recently, numerous cases of acute limb ischemia (ALI) have been reported although pooled data are lacking.

Methods: We systematically searched PubMed, Embase, Scopus, and the Cochrane Library for studies published online up to January 2021 that reported cases with SARS-CoV-2 infection and ALI. Eligible studies should have reported early outcomes including mortality. Primary endpoints included also pooled amputation, clinical improvement, and reoperation rates.

Results: In total, 34 studies (19 case reports and 15 case series/cohort studies) including a total of 540 patients (199 patients were eligible for analysis) were evaluated. All studies were published in 2020. Mean age of patients was 61.6 years (range, 39–84 years; data from 32 studies) and 78.4% of patients were of male gender (data from 32 studies). There was a low incidence of comorbidities: arterial hypertension, 49% (29 studies); diabetes mellitus, 29.6% (29 studies); dyslipidemia, 20.5% (27 studies); chronic obstructive pulmonary disease, 8.5% (26 studies); coronary disease, 8.3% (26 studies); and chronic renal disease, 7.6% (28 studies). Medical treatment was selected as first-line treatment for 41.8% of cases. Pooled mortality rate among 34 studies reached 31.4% (95% confidence interval [CI], 25.4%–37.7%). Pooled amputation rate among 34 studies reached 23.2% (95% CI, 17.3%–29.7%). Pooled clinical improvement rate among 28 studies reached 66.6% (95% CI, 55.4%–76.9%). Pooled reoperation rate among 29 studies reached 10.5% (95% CI, 5.7%–16.7%). Medical treatment was associated with a higher death risk compared with any intervention (odds ratio, 4.04; 95% CI, 1.075–15.197; $P = .045$) although amputation risk was not different between the two strategies (odds ratio, 0.977; 95% CI, 0.070–13.600; $P = .986$) (data from 31 studies).

Conclusions: SARS-CoV-2 infection is associated with a high risk for thrombotic complications, including ALI. COVID-associated ALI presents in patients with a low incidence of comorbidities, and it is associated with a high mortality and amputation risk. Conservative treatment seems to have a higher mortality risk compared with any intervention, although amputation risk is similar. (*J Vasc Surg* 2022;75:326–42.)

Keywords: Acute limb ischemia; COVID-19; SARS-CoV-2

The severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), the virus that causes coronavirus disease 2019 (COVID-19), originated in December 2019 and has caused a worldwide pandemic.¹ Infection with SARS-CoV-2 has been shown to have a wide range of clinical presentations from asymptomatic in a large percentage of

patients to devastating pulmonary failure, sepsis, and death.^{2,3} Additionally, the hypercoagulability associated with this infection has been recognized as a significant cause of morbidity, resulting in thrombotic complications such as pulmonary parenchymal thrombosis, venous thrombosis, myocardial infarction, and stroke.⁴ Lately, there have been reports of acute limb ischemia (ALI) observed among infected patients as well.

Although pooled data have been published on other thrombotic presentations such as acute mesenteric ischemia or stroke,^{5,6} pooled data on ALI are lacking. Therefore, aim of this review was to evaluate pooled data on patients with COVID-19 infection and ALI.

METHODS

Data sources and search. We systematically searched PubMed, Embase, Scopus, and the Cochrane Library (up to January 2021) for clinical studies published online that included patients suffered from ALI while diagnosed with COVID-19 infection. Eligible studies should have

From the Vascular Surgery Unit, First Department of Propedeutic Surgery, National and Kapodistrian University of Athens, Hippocraton Hospital^a; and the Department of Microbiology, National and Kapodistrian University of Athens, Areteion Hospital.^b

Author conflict of interest: none.

Correspondence: George Galyfos, MD, PhD, Vascular Surgery Unit, First Department of Propedeutic Surgery, National and Kapodistrian University of Athens, Hippocraton Hospital, 114 Vasilissis Sofias Ave, Athens, Greece 11527 (e-mail: georgegalyfos@hotmail.com).

The editors and reviewers of this article have no relevant financial relationships to disclose per the JVS policy that requires reviewers to decline review of any manuscript for which they may have a conflict of interest.

0741-5214

Copyright © 2021 by the Society for Vascular Surgery. Published by Elsevier Inc.

<https://doi.org/10.1016/j.jvs.2021.07.222>

reported at least early (30-day) mortality among other outcomes. This review was conducted according to established methods for systematic reviews in cardiovascular medicine (PRISMA criteria).⁷ The following medical subject terms were used for the online search: "acute," "limb," "ischemia," "COVID-19," "SARS-CoV-2," and "infection." In addition to searching databases, the reference lists of all included studies, meta-analyses, and reviews were evaluated manually, including unpublished data. Only studies published in English were included in this review. References from eligible articles or textbooks were also reviewed to identify further potential sources.

Data extraction: Outcomes and definitions. Three authors independently completed data extraction after following search criteria and quality assessment. Disagreements were resolved by consensus or after review by the senior author of the study, when necessary. Data were obtained from tables, graphs, and text as well. When data were presented in percentages, the absolute values were calculated. For each study, the following data were collected: first author, year of publication, country of origin, total number of patients included in the studies, total number of patients with ALI and COVID-19 infection, patient characteristics (mean age, gender, and comorbidities when reported), localization of ischemia (type of limbs, type of arteries), type of symptoms owing to infection (eg, fever, dyspnea, intubation need), type of medical treatment, type of interventional treatment, early outcomes (eg, mortality, amputation, and cardiac events), improvement of ischemia, late follow-up, and late outcomes when reported.

ALI was defined as acute ischemia presented in the lower or upper extremities. The cause of ischemia should have been arterial thrombosis or embolism and this cause should have been confirmed with some type of imaging within the included studies. Any cases with superficial or cutaneous limb necrosis in patients with COVID-19 infection who had no imaging evidence showing thrombosis or embolism of limb arteries were not included in this review. If the studies under evaluation included patients with acute ischemia of other type (such as mesenteric or cerebral), only patients with ALI were included in the analysis. If a patient happened to present with ALI and another type of ischemic complication, he or she was still included in the analysis.

Primary endpoints included early (30-day) mortality, amputation, clinical improvement, and reoperation rates.

Affected limb arteries included the subclavian, axillary, brachial, ulnar, radial, palmar, and digital arteries for the upper limb and the aorta, iliac, femoral, popliteal, tibial, peroneal, and plantar arteries for the lower limb.

All eligible patients should have tested positive for COVID-19 infection whether they presented with typical symptoms or not.

Amputations reported in this review included both major and minor amputations. Major amputations included transfemoral and transbrachial amputations as well as amputations below the level of the knee or elbow. Minor amputations included amputations below the level of the ankle or wrist. Pooled amputation rate included primary as well as secondary amputations.

Cardiac adverse events included cardiac arrest, myocardial infarction, arrhythmias, or acute cardiac failure.

Improvement of limb ischemia included the improvement of clinical symptoms/signs of ischemia without the need for further intervention or amputation.

Comorbidities are reported in the same way as reported in the included studies owing to lack of specific definitions in the majority of studies.

Quality assessment. Three authors independently reviewed study eligibility and quality. Disagreements were resolved by consensus or after review by the senior author of the study, when necessary. The quality of each study was assessed using well-established criteria for nonrandomized studies, specifically evaluating the collection of data, the aim of the studies, incomplete outcome data, statistical analysis, and other sources of bias.⁸ The quality of each study was evaluated and reported as high, medium, or low based on the design and methodology of study according to these criteria.

Inclusion and exclusion criteria. Studies included in this meta-analysis met the following criteria: (i) clinical studies or reports presenting cases with COVID-19 infection and ALI and (ii) studies should have reported early (30-day) mortality at least. ALI cases should have been documented with some type of imaging in the included studies. Studies reporting patients with acute ischemia at different body locations were also included but only patients with ALI from these studies were eligible for analysis.

Exclusion criteria included (i) types of publication other than clinical studies or reports, such as reviews, meta-analyses or editorials; (ii) studies not reporting at least early mortality among outcomes; (iii) studies presenting patients with superficial or cutaneous necrosis without evidence of an arterial thrombosis or embolism; (iv) studies reporting cases of ALI among patients without COVID-19 infection only; (v) studies reporting cases with COVID-19 infection and acute ischemia of other body parts such as mesenteric or cerebral ischemia; (vi) studies reporting cases with COVID-19 infection and acute thrombotic events without reporting outcomes for limb ischemia separately; (vii) studies published in a language other than English; (viii) studies not referring to humans; and (ix) studies reported as only abstracts or presented at conferences.

Statistical analysis. A meta-analysis was carried out using the StatsDirect Statistical software (Version 2.8.0,

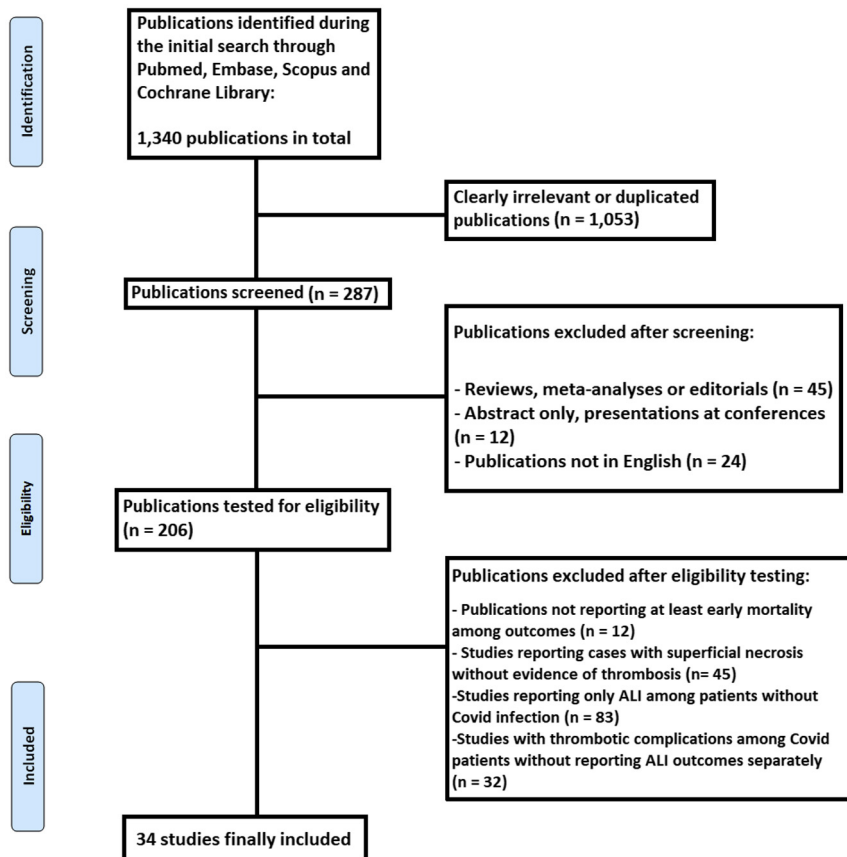


Fig 1. PRISMA flowchart of this review. ALI, Acute limb ischemia.

StatsDirect Ltd, Cambridge, UK). Odds ratios (OR) were used to determine effect size, along with 95% confidence intervals (CI). Regarding major outcomes, ORs were pooled with Der Simonian and Laird random effects models being used for sensitivity analysis. *P* values were calculated for evaluating statistical significance, with a *P* value of less than .05 indicating a statistically significant difference. Interstudy variations and heterogeneities were estimated using the *Q*-statistic with a *P* value of less than .05 also indicating a statistically significant heterogeneity. The present meta-analysis also quantified the effect of heterogeneity by using the *I*² index (range, 0%-100%), which represents the proportion of inter-study variability attributed to heterogeneity, rather than to chance.

In circumstances where more than one study reported data from the same cohort (introducing the potential for duplicate inclusion of patients), only the largest cohort was included in the main analysis. A χ^2 test with Yate's correction was used for comparing categorical variables between the two groups of patients. All statistical analyses were conducted using the absolute values and not percentages. The risk of bias was also assessed applying the Habbord-Egger test.

RESULTS

In total, 34 eligible studies (19 case reports and 15 case series/cohort studies) were included⁹⁻⁴² (Fig 1). Regarding quality, 5 studies were of high quality, 5 studies of medium quality, and 24 studies (mainly case reports) of lower quality. These studies evaluated a total of 540 patients, out of which 199 were eligible for this analysis. Table I presents basic characteristics of the included studies. The mean age of the patients was 61.6 years (range, 39-84 years; data from 32 studies) and 78.4% of patients were male (data from 32 studies), although data on age or gender were not provided by the two largest studies in size. Among 138 cases, the following limbs were affected: lower limb (n = 102), upper limb (n = 27), and bilateral lower limbs (n = 9). The exact affected arteries are presented in Table I as well.

Pooled demographics were the following: arterial hypertension, 49% (29 studies); diabetes mellitus, 29.6% (29 studies); dyslipidemia, 20.5% (27 studies); chronic obstructive pulmonary disease, 8.5% (26 studies); coronary disease, 8.3% (26 studies); arrhythmias, 14.1% (28 studies); chronic heart failure, 8.0% (25 studies); and chronic renal disease, 7.6% (28 studies). All comorbidities are presented in Table II.

Table I. Basic characteristics of the included studies

Study	Year of publication	Country of origin	No. of patients with COVID infection and ALI (total number of patients in the study)	Male/female gender	Mean age, years [SD or range are reported when provided by the studies]	Limb affected	Arteries affected
Veerasuri et al ⁹	2020	UK	1	1/0	56	Bilateral lower limbs	Right SFA, right trifurcation, left trifurcation
Kaur et al ¹⁰	2020	USA	1	1/0	43	Right lower limb	Right SFA, POPA, trifurcation
Brugliera et al ¹¹	2020	Italy	3	3/0	68	Bilateral lower limbs (n = 1) Right lower limb (n = 2)	SFA, POPA (n = 1) NR for 2 patients
Hanif et al ¹²	2020	Pakistan	1	0/1	75	Left upper limb	Left UA and RA
Hasan et al ¹³	2020	Pakistan	1	1/0	60	Right lower limb	Right POPA and trifurcation
Gubitosa et al ¹⁴	2020	USA	1	1/0	65	Right lower limb	Right POPA
Anwar et al ¹⁵	2020	USA	1	1/0	58	Left lower limb	Left SFA, trifurcation
Singh et al ¹⁶	2020	USA	1	1/0	77	Left lower limb	Left SFA, trifurcation
Wang et al ¹⁷	2020	USA	2	2/0	54	Right lower limb Right upper limb	Right dorsalis pedis, toes Right digital arteries, digits
Goldman et al ¹⁸	2020	USA	16 (48)	9/7	70 ± 14	Lower limbs (n = 16)	From the POPA and proximally (n = 15) Distally from the POPA (n = 1)
Sánchez et al ¹⁹	2020	Peru	30	23/7	60 ± 15	Lower limb (n = 22) Upper limb (n = 8)	NR
Galanis et al ²⁰	2020	Greece	1	1/0	80	Right Upper limb	RA and UA
Bozzani et al ²¹	2020	Italy	6 (38)	4/2	71 (49-83)	Lower limb (n = 6)	Iliac-SFA-POPA (n = 6)
Mietto et al ²²	2020	Italy	1	1/0	53	Left lower limb	Iliac-SFA-POPA-tibial arteries
Baccellieri et al ²³	2020	Italy	1	1/0	67	Right lower limb	Iliac-SFA-POPA
Shao et al ²⁴	2020	USA	1	1/0	67	Right upper limb	Brachial artery, UA, RA
Etkin et al ²⁵	2020	USA	42 (49)	NR	NR	Lower extremities (n = 35) Upper limbs (n = 7)	Aortoiliac (n = 8) Femoral (n = 12) POPA (n = 15) Above elbow (n = 4) UA and RA (n = 3)
Muhammad et al ²⁶	2020	UK	1	1/0	49	Left lower limb	Aorti-iliac-POPA-trifurcation

(Continued on next page)

Table I. Continued.

Study	Year of publication	Country of origin	No. of patients with COVID infection and ALI (total number of patients in the study)	Male/female gender	Mean age, years [SD or range are reported when provided by the studies]	Limb affected	Arteries affected
Heald et al ²⁷	2020	USA	1	1/0	65	Left hand-digital ischemia	Digital arteries (left first and second digits)
Bellosta et al ²⁸	2020	Italy	20	18/2	75 ± 8	Upper and lower limbs (number NR)	NR
Kaur et al ²⁹	2020	USA	1	1/0	71	Upper limb (right)	Right brachial artery and RA
Kahlberg et al ³⁰	2020	Italy	41 (305)	NR	NR	Upper and lower limbs	NR
Schultz et al ³¹	2020	USA	2	1/1	57	3 fingers (right) 2 fingers (right)	Digital arteries (right) RA (right)
Vacirca et al ³²	2020	Italy	1	1/0	58	Right lower limb	ATA, PTA, PA (right)
Fan et al ³³	2020	Singapore	1	1/0	39	Right lower limb	ATA Abdominal aorta
Perini et al ³⁴	2020	Italy	2	2/0	45	Bilateral lower limbs Upper limb (left)	Aortoiliac Brachial bifurcation
Kashi et al ³⁵	2020	France	5 (7)	4/1	69	Lower limbs (2 bilateral, 2 right, 1 left)	Femoral (n = 3) POPA (n = 1) Iliac (n = 1) Fem-pop bypass (n = 1) NR (n = 1)
Baeza et al ³⁶	2020	Spain	3	1/2	72	Bilateral lower limbs (n = 3)	Aortic-bilateral iliac (n = 3)
Garg et al ³⁷	2020	USA	3 (4)	2/1	63	Lower limb (2 right, 1 left)	POPA (n = 3)
Thompson et al ³⁸	2020	USA	1	0/1	42	Right upper limb	Right subclavian + UA
Levolger et al ³⁹	2020	Netherlands	2 (4)	2/0	53	Right lower limb Left upper limb	Right common iliac artery Left subclavian artery
Wengerter et al ⁴⁰	2020	USA	3 (4)	3/0	53	Left lower limb (n = 2) Bilateral lower limbs (n = 1)	Aortoiliac (n = 1) Femoral-POPA (n = 3)
Chowdhury et al ⁴¹	2020	USA	1	1/0	75	Right upper extremity	Brachial artery
Liu et al ⁴²	2021	China	1	1/0	70	Right lower limb	CFA and SFA

ATA, Anterior tibial artery; CFA, common femoral artery; NR, not reported; POPA, popliteal artery; RA, radial artery; PA, peroneal artery; PTA, posterior tibial artery; SD, standard deviation; SFA, superficial femoral artery; UA, ulnar artery.

Concerning COVID-19 infection, 49.1% of patients presented with fever (27 studies), 62.3% of patients presented with dyspnea (27 studies), and 36.4% of patients needed to be intubated (30 studies). Basic laboratory findings are also presented in Table III. Increased D-Dimers levels (>5 µg/mL; 26 studies) were not associated with

death (OR, 1.169; 95% CI, 0.360-3.756; $P = .792$) or amputation (OR, 2.0; 95% CI, 0.334-11.969; $P = .448$) risk. Increased C-reactive protein (CRP) levels (>20 mg/L; 16 studies) were not associated with death (OR, 3.261; 95% CI, 0.164-65.012; $P = .438$) or amputation (OR, 3.627; 95% CI, 0.183-72.070; $P = .398$) risk. Increased fibrinogen levels

Table II. Demographics of the included patients

Study	Arterial hypertension	Diabetes mellitus	Dyslipidemia	COPD	CAD	Arrhythmia	CHF	Renal Disease	Other comorbidities
Veerasuri et al ⁹	0	0	0	0	0	0	0	0	None
Kaur et al ¹⁰	1/1	1/1	0	0	0	0	0	0	None
Brugliera et al ¹¹	3/3	2/3	1/3	1/3	0	0	0	1/3	Hypothyroidism (n = 1)
Hanif et al ¹²	0	0	0	0	0	0	0	0	None
Hasan et al ¹³	1/1	0	0	0	0	0	0	0	None
Gubitosa et al ¹⁴	1/1	1/1	1/1	0	0	0	0	1/1	Smoking history
Anwar et al ¹⁵	0	0	0	0	0	0	0	0	None
Singh et al ¹⁶	0	0	0	0	0	0	0	0	None
Wang et al ¹⁷	0	0	0	0	0	0	0	0	None
Goldman et al ¹⁸	13/16	8/16	8/16	NR	NR	NR	4/16	0/16	Smoking history (n = 8) PAD (n = 8)
Sánchez et al ¹⁹	10/30	8/30	1/30	NR	4/30	3/30	NR	1/30	Smoking history (n = 3) PAD (n = 4)
Galanis et al ²⁰	1/1	1/1	0	0	0	0	0	0	Dementia
Bozzani et al ²¹	NR	NR	NR	NR	NR	NR	NR	NR	NR
Mietto et al ²²	1/1	0	0	0	0	0	0	0	Obesity
Baccellieri et al ²³	0	0	0	0	0	0	0	0	Obesity
Shao et al ²⁴	0	0	0	0	0	0	0	0	Lupus anticoagulant positive
Etkin et al ²⁵	NR	NR	NR	NR	NR	NR	NR	NR	NR
Muhammad et al ²⁶	NR	NR	NR	NR	NR	NR	NR	NR	NR
Heald et al ²⁷	1/1	0	0	0	0	0	0	0	History of smoking
Bellosta et al ²⁸	11/20	3/20	NR	2/20	2/20	5/20	NR	4/20	Previous VS (n = 4) Obesity (n = 4)
Kaur et al ²⁹	0	1/1	0	0	0	0	0	0	None
Kahlberg et al ³⁰	NR	NR	NR	NR	NR	NR	NR	NR	NR
Schultz et al ³¹	1/2	0	1/2	0	0	0	0	0	Obesity (n = 1)
Vacirca et al ³²	NR	NR	NR	NR	NR	NR	NR	NR	NR
Fan et al ³³	0	0	0	0	0	0	0	0	None
Perini et al ³⁴	0	0	0	0	0	0	0	0	None
Kashi et al ³⁵	4/5	1/5	NR	1/5	NR	2/5	NR	1/5	Smoking (n = 2) PAD (n = 2)
Baeza et al ³⁶	2/3	1/3	2/3	1/3	0	2/3	0	0	Smoking (n = 2) Obesity (n = 1) PAD (n = 1)
Garg et al ³⁷	1/3	1/3	1/3	NR	NR	1/3	NR	NR	NR
Thompson et al ³⁸	0	1/1	0	0	0	0	0	0	Rheumatoid arthritis
Levolger et al ³⁹	0	1/2	0	0	0	0	0	0	-
Wengerter et al ⁴⁰	1/3	2/3	1/3	0	0	0	0	0	Smoking (n = 1)
Chowdhury et al ⁴¹	1/1	0	1/1	0	1/1	0	0	0	Dementia
Liu et al ⁴²	0	0	0	0	0	0	0	0	Lung cancer

CAD, Coronary artery disease; CHF, chronic heart failure; COPD, chronic obstructive pulmonary disease; NR, not reported; PAD, peripheral artery disease; VS, vascular surgery.

Table III. Symptoms and laboratory findings of the included patients

Study	Fever	Dyspnea	Need for intubation	Mean leucocyte count ($\times 10^9/L$)	Mean platelet count ($\times 10^9/L$)	Fibrinogen (mg/dL)	CRP (mg/L)	D-Dimers ($\mu g/mL$)	PCT (ng/mL)
Veerasuri et al ⁹	1/1	1/1	0	0.8	NR	NR	NR	23.138	NR
Kaur et al ¹⁰	1/1	1/1	1/1	16	484	853	289.7	20	67
Brugliera et al ¹¹	1/3	3/3	1/3	NR	Normal for all	466.3	NR	20, NR, 5.04	NR
Hanif et al ¹²	1/1	0	0	5.9	733	NR	NR	0.867	NR
Hasan et al ¹³	1/1	1/1	0	13.9	95	NR	82.5	1.96	0.3
Gubitosa et al ¹⁴	1/1	1/1	1/1	Lymphopenia	82	247	NR	7.955	NR
Anwar et al ¹⁵	1/1	1/1	0	NR	NR	312	25.23	15.653	25.23
Singh et al ¹⁶	0	1/1	0	41	534	NR	301	2.77	0.6
Wang et al ¹⁷	2/2	2/2	2/2	NR	NR	NR	NR	8.25	NR
Goldman et al ¹⁸	3/16	8/16	4/16	13.5 \pm 4	NR	NR	NR	NR	NR
Sánchez et al ¹⁹	NR	NR	NR	11.6 [9.7-16.1]	284 [220-371]	4.8 [4.7-6.3]	35.5 [24-61]	3.2 [1.6-4.3]	NR
Galanis et al ²⁰	1/1	1/1	1/1	5.6	174	360	166	13.6	0.1
Bozzani et al ²¹	NR	NR	2/6	NR	NR	NR	78.17 [3-240]	NR	NR
Mietto et al ²²	0	0	0	NR	NR	NR	NR	NR	NR
Baccellieri et al ²³	1/1	1/1	0	NR	NR	711	114.1	20	NR
Shao et al ²⁴	0	1/1	1/1	Increased	NR	NR	NR	>5	NR
Etkin et al ²⁵	NR	NR	NR	NR	NR	NR	NR	NR	NR
Muhammad et al ²⁶	1/1	1/1	0	11.8	520	NR	12	NR	NR
Heald et al ²⁷	0	1/1	1/1	NR	NR	NR	NR	0.79	NR
Bellosta et al ²⁸	NR	NR	NR	14 \pm 2	239 \pm 82	NR	NR	2.2	NR
Kaur et al ²⁹	1/1	1/1		8.6	331	-	111.9	1.85	-
Kahlberg et al ³⁰	NR	NR	NR	NR	NR	NR	NR	NR	NR
Schultz et al ³¹	1/2	2/2	2/2	NR	NR	486, NR	25, NR	7.56	NR
Vacirca et al ³²	0	1/1	1/1	6.07	322	524	NR	1.19	0.1
Fan et al ³³	1/1	1/1		NR	NR	770	136.2	2.55	NR
Perini et al ³⁴	NR	NR	1/2	NR	NR	NR	NR	9.0	NR
Kashi et al ³⁵	NR	NR	3/5	NR	160	NR (n = 2) 547	NR	NR (n = 3) 20	NR
Baeza et al ³⁶	3/3	0/3	0/3	18.2	193.7	766.3	4.1	5.07	NR
Garg et al ³⁷	1/3	2/3	1/3	NR	NR	NR	NR	3.3	NR
Thompson et al ³⁸	0	0	0	NR	NR	NR	NR	NR	NR
Levolger et al ³⁹	2/2	1/2	0/2	10.6	360	NR	167	NR	NR
Wengerter et al ⁴⁰	1/3	0	1/3	NR	NR	NR	144.3	15.6	NR
Chowdhury et al ⁴¹	1/1	1/1	1/1	8.6	172	NR	20	1.2	NR
Liu et al ⁴²	0	0	0	9.83	282	560	79.1	6.55	NR

CRP, C-reactive protein; NR, not reported; PCT, procalcitonin. Laboratory values are reported either with standard deviation or value range, whatever was reported.

(>400 mg/dL; 13 studies) were not associated with death (OR, 0.667; 95% CI, 0.152-2.925; $P = .485$) or amputation (OR, 0.639; 95% CI, 0.184-2.222; $P = .481$) risk.

Regarding treatment, medical treatment was chosen in 41.8% of patients (33 studies) as a first-line treatment. In total (among 33 studies), 92 patients (58.2%) underwent the following procedures: thrombectomy (n = 81), fasciotomy (n = 9), angioplasty with or without stenting

(n = 7), thrombolysis (n = 7), thrombosuction (n = 2), bypass (n = 3), and endarterectomy (n = 2). All patients were covered with unfractionated or low-molecular-weight heparin. All medical and interventional treatment is presented in Table IV.

The pooled mortality rate among 34 studies reached 31.4% (95% CI, 25.4%-37.7%). The pooled amputation rate (both primary and secondary) among 34 studies reached

Table IV. Type of treatment for the included patients

Study	Medical to interventional treatment as first line treatment	HCQ	Antibiotics	Antiviral treatment	Heparin or LMWH	Other medication	Thrombectomy or Embolectomy	Other intervention	Reoperation
Veerasuri et al ⁹	1/0	NR	NR	NR	1/1	Rivaroxaban	0	None	0
Kaur et al ¹⁰	1/0	1/1	1/1	NR	1/1	NR	0	None	0
Brugliera et al ¹¹	2/1	3/3	2/3	1/3	3/3	Iloprost (n = 1) ASA (n = 2)	1	None	0
Hanif et al ¹²	0/1	0	1/1	0	1/1	NR	1	None	0
Hasan et al ¹³	1/0	0	1/1	0	1/1	Rivaroxaban post discharge	0	None	0
Cubitosa et al ¹⁴	0/1	0	0	0	1/1	FDPX	1	Fasciotomy	0
Anwar et al ¹⁵	0/1	1/1	1/1	0	1/1	CTDS, ASA	0	Angioplasty	0
Singh et al ¹⁶	0/1	1/1	1/1	0	1/1	NR	1	None	0
Wang et al ¹⁷	2/0	NR	NR	NR	1/1	Argatroban	0	None	0
Goldman et al ¹⁸	9/7	NR	NR	NR	NR	NR	6	None	0
Sánchez et al ¹⁹	2/28	NR	NR	NR	NR	NR	23	Fasciotomy (n = 6)	2
Galanis et al ²⁰	1/0	1/1	1/1	0	1/1	FDPX	1	None	0
Bozzani et al ²¹	0/6	NR	NR	NR	6/6	ASA (3/6) ASA + clopidogrel (3/6)	6	PTA (n = 2) PTA + stenting (n = 1)	1
Mietto et al ²²	0/1	NR	NR	NR	1/1	Prostacyclin	1	Thrombolysis, fasciotomy	1
Baccellieri et al ²³	0/1	1/1	1/1	NR	1/1	NR	1	Thrombectomy at right upper limb	0
Shao et al ²⁴	0/1	NR	NR	NR	1/1	NR	1	Thrombolysis, fasciotomy	0
Etkin et al ²⁵	31/11	NR	NR	NR	NR	NR	9	Endovascular thrombosuction (n = 2)	NR
Muhammad et al ²⁶	0/1	NR	NR	NR	1/1	ASA 75 mg Dabigatran 150 mg bid	0	Thrombolysis	0
Heald et al ²⁷	1/0	NR	NR	NR	1/1	NR	0	None	0
Bellosta et al ²⁸	3/17	NR	NR	NR	20/20	NR	15	Below the knee fempop bypass (n = 2) Additional thrombolysis (n = 2) Kissing stents (n = 2) Femoral endarterectomy (n = 1) Below the knee angioplasty (n = 1)	2
Kaur et al ²⁹	0/1	1/1	1/1	NR	1/1	NR	1	Endarterectomy of the right arm	0
Kahlberg et al ³⁰	NR	NR	NR	NR	NR	NR	NR	NR	NR
Schultz et al ³¹	2/0	2/2	2/2	2/2	2/2	Nitroglycerin (n = 2) Apixaban (n = 1)	0	None	0
Vacirca et al ³²	0/1	NR	NR	NR	1/1	NR	1	Thrombolysis	0
Fan et al ³³	0/1	NR	NR	1/1	1/1	ASA	1	Aortic stent graft placement	0
Perini et al ³⁴	1/1	NR	NR	NR	1/1	NR	1	None	1
Kashi et al ³⁵	4/1	NR	NR	NR	2/5	Apixaban (n = 1) ASA (n = 2)	1	None	NR
Baeza et al ³⁶	0/3	3/3	3/3	2/3	3/3	Acenocoumarol (n = 1)	2	Aortobifemoral bypass (n = 1)	0
Garg et al ³⁷	2/1	1/3	1/3	1/3	3/3	–	1	NR	NR
Thompson et al ³⁸	0/1	0	0	0	1/1	–	1	None	0
Levolger et al ³⁹	2/0	1/2	0	0	2/2	Rivaroxaban (n = 1) Apixaban (n = 1)	1	Thrombolysis	0
Wengert et al ⁴⁰	0/3	NR	NR	NR	3/3	NR	3	None	NR

(Continued on next page)

Table IV. Continued.

Study	Medical to interventional treatment as first line treatment	HCQ	Antibiotics	Antiviral treatment	Heparin or LMWH	Other medication	Thrombectomy or Embolectomy	Other intervention	Reoperation
Chowdhury et al ⁴¹	0/1	1/1	1/1	0	1/1	Prednisolone	1	None	0
Liu et al ⁴²	1/0	NR	1/1	1/1	1/1	NR	0	None	0
Total (n/total n)	66/158 (1 study NR)	17/24 (18 studies NR)	18/24 (18 studies NR)	8/23 (19 studies NR)	65/65 (4 studies NR)	ASA (n = 13) Clopidogrel (n = 3) Apixaban (n = 3) Dabigatran (n = 1) Argatroban (n = 1) Rivaroxaban (n = 3) CTDS (n = 2) FDPX (n = 1) Acenocoumarol (n = 1) Prostaglandins (n = 2) Nitroglycerine (n = 2) (15 studies NR)	N = 81 (1 study NR)	Thrombolysis (n = 7) PTA/stenting (n = 5) Endarterectomy (n = 2) Fasciotomy (n = 9) Bypass (n = 3) Thrombectomy at other site (n = 1) Thrombosuction Stent graft placement (n = 1)	n = 7 (5 studies NR)

ASA, Acetylsalicylic acid; *bid*, 2 times per day; CTDS, corticosteroids; FDPX, fondaparinux; HCQ, hydroxychloroquine; LMWH, low molecular weight heparin; NR, not reported; PTA, percutaneous transluminal angioplasty.

23.2% (95% CI, 17.3%-29.7%). From the 34 reported amputations, 22 were major (transfemoral or below the knee), 1 minor (below the ankle), and 11 were at an unknown level. The pooled clinical improvement rate among 28 studies reached 66.6% (95% CI, 55.4%-76.9%). The pooled reoperation rate among 29 studies reached 10.5% (95% CI, 5.7%-16.7%) (Figs 2-5). For all these pooled outcomes, heterogeneity was very low ($I^2 = 0\%-7.7\%$). All outcomes are presented in Table V. Finally, there was no difference regarding death risk (OR, 1.10; 95% CI, 0.284-4.272; $P = .884$) and amputation risk (OR, 1.16; 95% CI, .288-4.649; $P = .834$) between upper limb and lower limb location (data available from 31 studies). Regarding first-line strategy, medical treatment was associated with a higher risk of death compared with any intervention (OR, 4.04; 95% CI, 1.075-15.197; $P = .045$), although the risk of amputation was not different between the two strategies (OR, 0.977; 95% CI, 0.070-13.600; $P = .986$) (data from 31 studies).

DISCUSSION

Although the overall incidence of ALI has decreased worldwide and the hypercoagulable state remains an uncommon cause for limb ischemia,⁴³ the incidence of thromboembolic complications among patients with COVID-19 infection is as high as 35% to 45%.⁴⁴ In critically ill patients, there is an even higher risk for both venous and arterial thromboembolism associated with high mortality.^{45,46} As we found in this review, ALI was associated with a mortality rate of 31.4%, although the reported mortality in non-COVID populations with ALI ranges from 5% to 9% in literature.^{47,48} Comparative studies have also shown a higher incidence of thrombotic events such as strokes among COVID-infected patients compared with the general wards.⁴⁹ However, this relative increase of arterial thrombotic events during the pandemic may be attributed to several factors, such as delays in

emergency room presentation owing to the lockdown, older patient age, or fear in approaching hospitals because of a high contamination risk.⁵⁰ Additionally, there are several reports that these thrombotic events occur at a later time point during the course of the infection.³⁶ Some authors have advocated that the virus starts a second attack between 7 and 14 days from the onset of symptoms that perhaps initiates some type of hypercoagulability.²

Additionally, we found a mean age of 61 years with a very wide age range starting from just 39 years. Other authors have also reported that infected patients with thrombotic complications are of relatively young age, and available computed tomography scans and angiography reveal no prior major atherosclerosis in these cases.⁵¹ This finding suggests that a significant proportion of arterial thromboses in patients with COVID-19 might occur over nondiseased or mildly diseased vessels. Although male gender, advanced age, hypertension, and diabetes have been found to be independent risk factors of death among patients with COVID-19,⁵² this review revealed that infected patients with ALI show a low incidence of major comorbidities such as diabetes, dyslipidemia, coronary disease, and renal disease. This finding indicates that even patients without risk factors are at risk of presenting thrombotic complications when infected.

The causative mechanism for ALI seems to be a systematic inflammatory process triggered by a massive activation of macrophages that generate a cytokine storm.⁵³ COVID-19 causes elevated cytokine levels, including but not limited to tumor necrosis factor- α , IL-1 β , IL-6, procalcitonin, and interferon γ .⁵⁴⁻⁵⁶ The coupling of inflammation and coagulation has also been described in the literature, with these procedures sharing common molecular pathways.⁵⁷ It has been reported that infected patients are prone to thrombotic dysfunction, and especially

Early mortality

Non-combinability of studies

Cochran Q = 31.50455 (df = 33)

P = 0.5416

Moment-based estimate of between studies variance = 0

I^2 (inconsistency) = 0% (95% CI = 0% to 34.9%)

Random effects (DerSimonian-Laird)

Pooled proportion = 0.31356 (95% CI = 0.253528 to 0.376903)

Bias indicators

Egger: bias = -0.199722 (95% CI = -1.563764 to 1.16432) P = 0.7674

Harbord: bias = -0.22576 (92.5% CI = -0.726201 to 0.27468) P = 0.4126

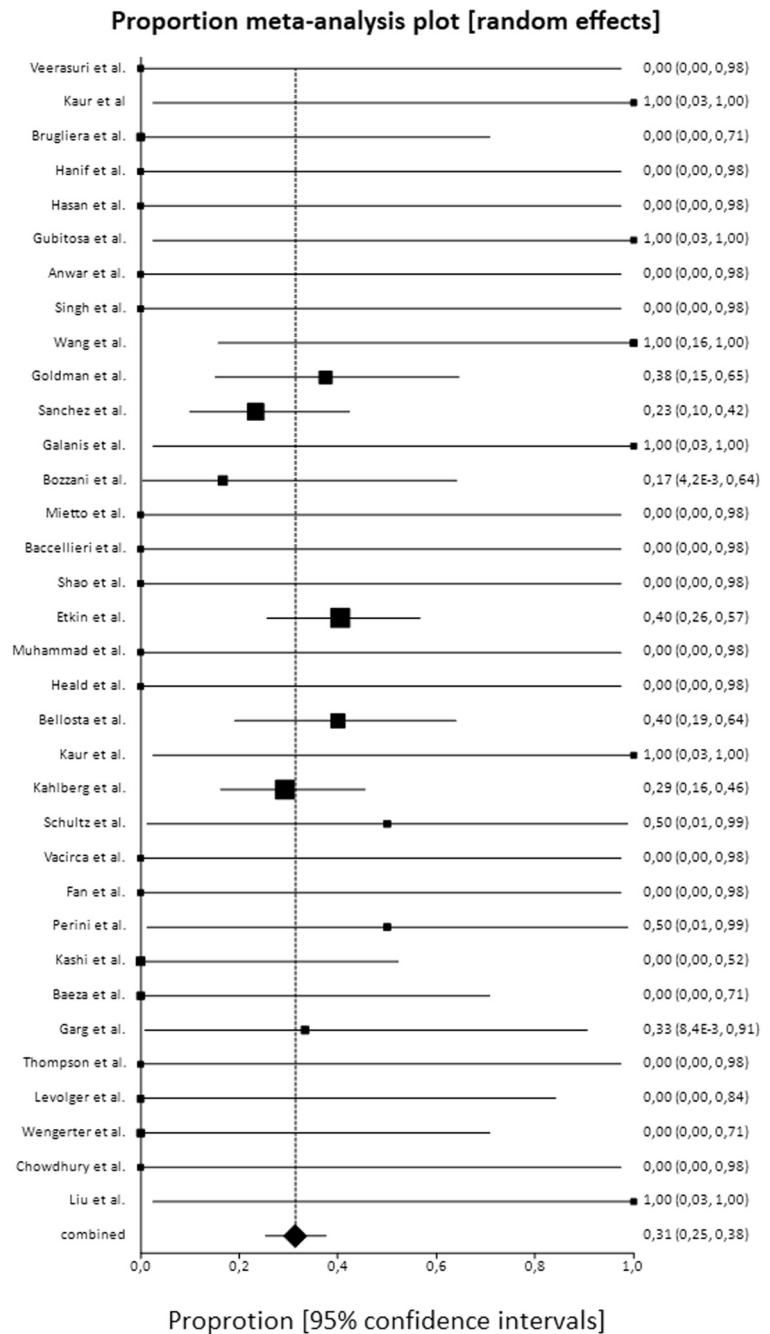


Fig 2. Forest plot on pooled early mortality. *CI*, Confidence interval; *df*, degrees of freedom.

those with severe symptoms had higher CRP levels and a higher thrombotic risk.⁵⁸ However, high levels of CRP in our review were not associated with adverse events in patients with ALI. These patients exhibit several risk factors of thrombosis such as blood concentration, vascular endothelial injury, extended bed rest, and blood hypercoagulation. Additionally, recent data indicate that COVID-19 infection is associated with profound and generalized activation of both alternative and lectin-based complement pathways.⁵⁹

There is growing evidence that this virus promotes a procoagulant state producing both microthrombi and macrothrombi.⁶⁰ Cutaneous ischemic lesions are frequent in such patients, even in absence of major vessel thrombosis.⁶¹ In particular, severe endothelial injury, widespread thrombosis with microangiopathy, alveolar-capillary microthrombi and new vessel growth have been detected in infected cases.⁶² Vascular pathological changes in such patients include partial vascular endothelial shedding, vascular intimal

Amputation rate

Non-combinability of studies

Cochran Q = 31.025712 (df = 32)

P = 0.5157

Moment-based estimate of between studies variance = 0

I^2 (inconsistency) = 0% (95% CI = 0% to 35.3%)

Random effects (DerSimonian-Laird)

Pooled proportion = 0.23197 (95% CI = 0.172518 to 0.297311)

Bias indicators

Egger: bias = 0.267629 (95% CI = -0.958615 to 1.493874) P = 0.6593

Harbord: bias = -0.062513 (92.5% CI = -0.63102 to 0.505994) P = 0.8408

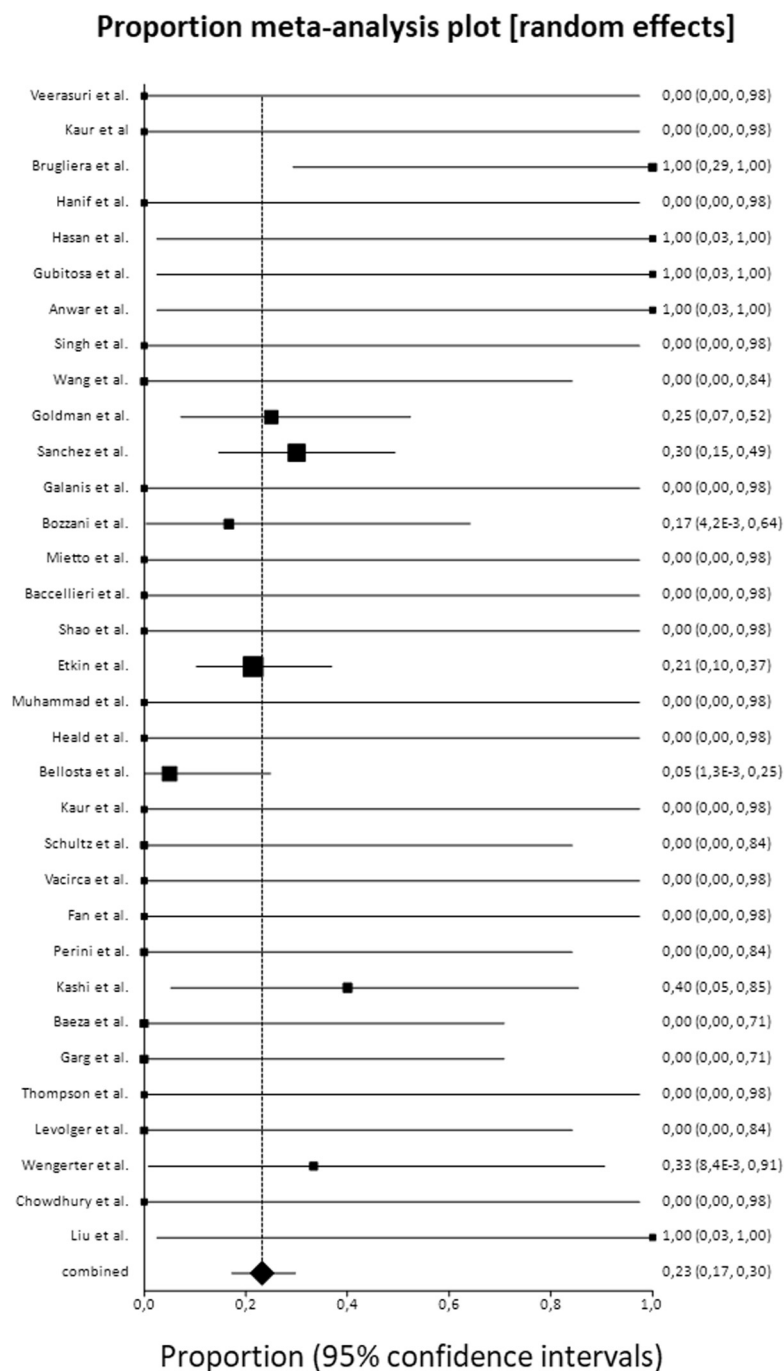


Fig 3. Forest plot on pooled amputation rate. *CI*, Confidence interval; *df*, degrees of freedom.

inflammation, and thrombosis.⁶³ Varga et al⁶³ have even observed viral inclusion bodies under light microscopy in the endothelium of the specimens. The angiotensin-converting enzyme 2, the receptor for SARS-CoV-2, is also expressed on the membrane of vascular muscle and endothelial cells, and therefore, infection of these cells could induce an inflammatory response in the blood vessel walls, predisposing to

clot formation.³³ The viral infection itself leads to decreased platelet function secondary to decreased production, platelet consumption, and production of autoantibodies such as antiphospholipid antibodies.^{64,65} It is also known that other viral infections including hepatitis or human immune deficiency viral infections have been associated with thrombotic complications such as venous thromboembolism.^{66,67}

Clinical improvement rate

Non-combinability of studies

Cochran Q = 29.265959 (df = 27)

P = 0.3481

Moment-based estimate of between studies variance = 0.028367

I₂ (inconsistency) = 7.7% (95% CI = 0% to 42.2%)

Random effects (DerSimonian-Laird)

Pooled proportion = 0.665777 (95% CI = 0.554272 to 0.768612)

Bias indicators

Egger: bias = -1.539691 (95% CI = -3.25217 to 0.172789) P = 0.076

Harbord: bias = -0.748902 (92.5% CI = -1.438007 to -0.059798) P = 0.0543

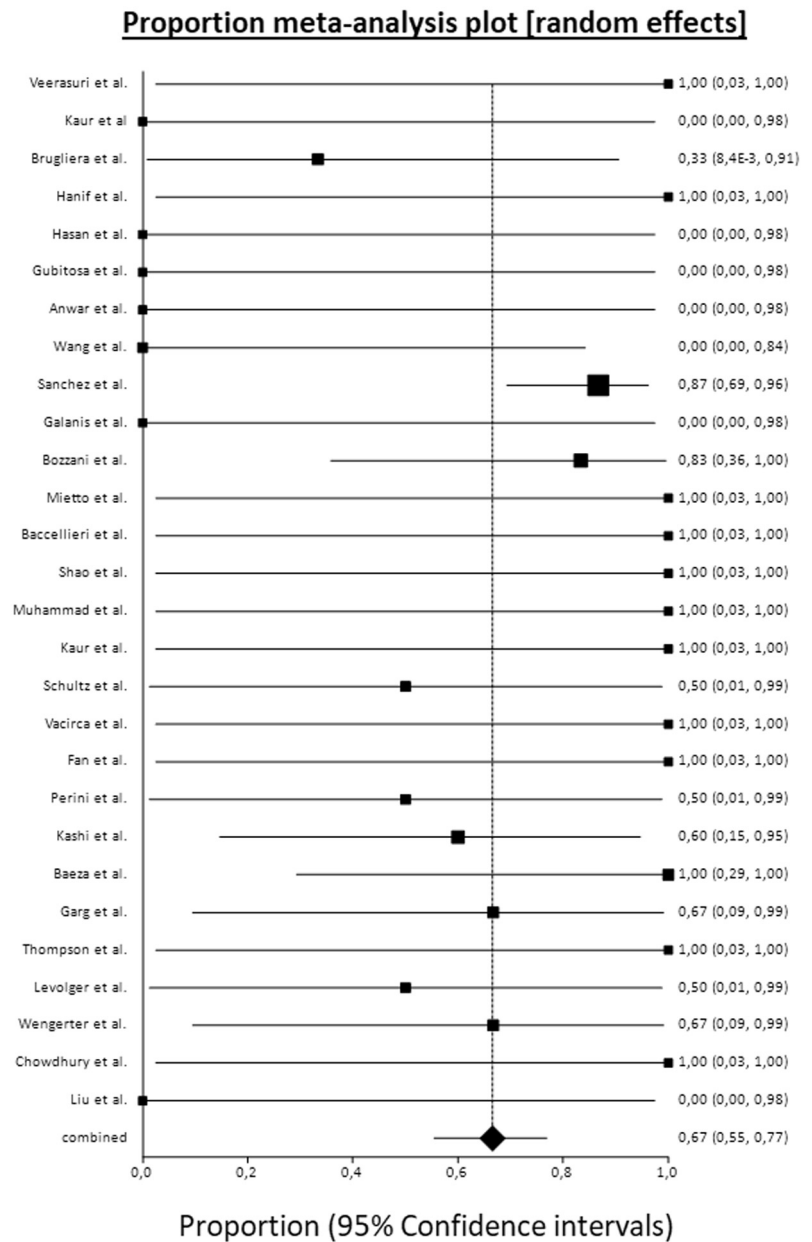


Fig 4. Forest plot on pooled clinical improvement rate. *CI*, Confidence interval; *df*, degrees of freedom.

Regarding thrombotic markers, high D-dimers, fibrinogen degradation products, and a prolonged thromboplastin time have been associated with greater in-hospital mortality, need for mechanical ventilation, and thrombotic complications in infected patients.^{12,68,69} Recent pooled data indicate that prothrombin time and D-dimer levels are significantly higher in patients with severe infection than in those with mild disease.⁷⁰ Some authors advocate that increased levels of D-dimers could serve as an indicator of the time-point at which an

intervention with recombinant tissue plasminogen activator or tocilizumab should be considered.⁷¹ However, an optimal cut-off level and prognostic value are still not known.⁷² In our review, all patients had a thrombotic complication, indicating a population at higher mortality risk. This factor could justify that we could not establish a cutoff level for D-dimer as well. The mean fibrinogen concentrations in patients with COVID-19 are in general at the upper limits of normal, presumably as an acute phase response.⁴⁴ However, we could not establish an

Reoperation rate

Non-combinability of studies

Cochran Q = 10.61848 (df = 28)

P = 0.9988

Moment-based estimate of between studies variance = 0

I^2 (inconsistency) = 0% (95% CI = 0% to 37%)

Random effects (DerSimonian-Laird)

Pooled proportion = 0.105424 (95% CI = 0.056983 to 0.166515)

Bias indicators

Egger: bias = 0.127973 (95% CI = -0.449968 to 0.705913) P = 0.6532

Harbord: bias = -0.012953 (92.5% CI = -0.464592 to 0.438686) P = 0.958

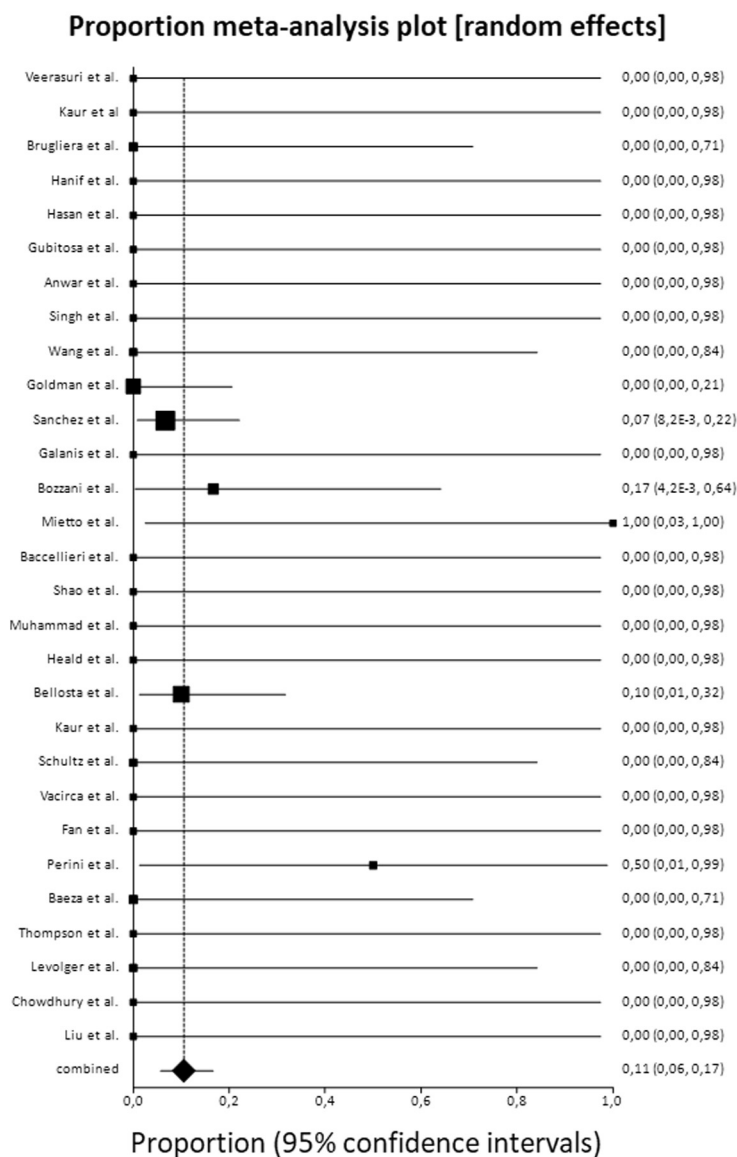


Fig 5. Forest plot on pooled reoperation rate. *CI*, Confidence interval; *df*, degrees of freedom.

association of high fibrinogen levels with worse outcomes either.

Given the increased thrombotic risk, the World Health Organization recommends at least prophylactic doses of low-molecular-weight heparin daily or subcutaneous unfractionated heparin twice daily for venous thromboembolism prophylaxis in critically ill patients with COVID-19.⁷³ However, ALI can even occur among patients already receiving thromboprophylaxis,⁷⁴ and this outcome has been observed in some cases included in the present review. Despite thromboprophylaxis, the risk of venous thromboembolism remains high in hospitalized patients with COVID-19.⁷⁵ One study of 94 patients with confirmed COVID-19 demonstrated a

statistically significant relative deficiency of antithrombin III compared with control.⁷⁶ This acquired deficiency would promote further coagulation and could decrease the efficacy of anticoagulant treatment in such patients. Therefore, certain concerns arise whether full therapeutic dosage of anticoagulants would be more appropriate for severely ill patients.

For patients with COVID-19 presenting with ALI, the choice of intervention is guided by the need to limit interventions that would expose these patients to stressful procedures, the desire to limit exposure of medical personnel, and the need to conserve resources. Additionally, in critically ill infected patients, thrombosis may be a terminal event, sometimes being referred to as agonal

Table V. Main outcomes for all included patients

Study	Death	Amputations	Cardiac	Other complications	Improvement of the ischemia	Mean follow-up	Late outcomes
Veerasuri et al ⁹	0	0	0	0	1	2 months	Numbness in right foot
Kaur et al ¹⁰	1	0	1	Hemodialysis	0	NR	NR
Brugliera et al ¹¹	0	3 (all TF)	0	DVT (n = 1)	1	NR	NR
Hanif et al ¹²	0	0	0	0	1	NR	NR
Hasan et al ¹³	0	1 (TF)	0	0	0	NR	NR
Cubitosa et al ¹⁴	1	1 (TF)	1	HIT	0	NR	NR
Anwar et al ¹⁵	0	1 (below ankle)	0	0	0	NR	NR
Singh et al ¹⁶	0	0	0	0	NR	NR	NR
Wang et al ¹⁷	2	0	0	Encephalopathy (n = 1)	0	NR	NR
Goldman et al ¹⁸	6	4 (all TF)	NR	NR	NR	NR	NR
Sánchez et al ¹⁹	7	9 (all TF)	NR	NR	26	NR	NR
Galanis et al ²⁰	1	0	0	DVT	0	NR	NR
Bozzani et al ²¹	1	1 (TF)	0	Rethrombosis (n = 2)	5	NR	NR
Mietto et al ²²	0	0	0	Rhabdomyolysis, acute renal failure	1	40 days	Superficial peroneal nerve impairment
Baccellieri et al ²³	0	0	0	Nephrotic syndrome, bilateral acroischemia, thrombosis of upper limb (intraoperative)	1	2 months	No new event
Shao et al ²⁴	0	0	0	PE, GI bleeding	1	2 months	Dry gangrene of digits
Etkin et al ²⁵	17	9 (NR)	NR	NR	NR	NR	NR
Muhammad et al ²⁶	0	0	0	NR	1	6 weeks	Full recovery
Heald et al ²⁷	0	0	0	0	NR	NR	NR
Bellosta et al ²⁸	8	1 (major)	1	NR	NR	NR	NR
Kaur et al ²⁹	1	0	1	NR	1	NR	NR
Kahlberg et al ³⁰	12	NR	NR	NR	NR	NR	NR
Schultz et al ³¹	1	0	0	Septic shock, ARDS, AKI (n = 2) DVT (n = 2) Hemorrhagic shock (n = 1)	1	NR	NR
Vacirca et al ³²	0	0	0	0	1	NR	NR
Fan et al ³³	0	0	0	NR	1	NR	NR
Perini et al ³⁴	1	0	0	0	1	NR	NR
Kashi et al ³⁵	0	2 (NR)	NR	DVT (n = 1)	3	NR	NR
Baeza et al ³⁶	0	0	0/3	0	3	NR	NR
Garg et al ³⁷	1	0	0	DVT (n = 1) Stroke (n = 1)	2	NR	NR
Thompson et al ³⁸	0	0	0	–	1	Several days	Digit tip gangrene
Levolger et al ³⁹	0	0	0	Contralateral limb ischemia (n = 1)	1	NR	NR

(Continued on next page)

Table V. Continued.

Study	Death	Amputations	Cardiac	Other complications	Improvement of the ischemia	Mean follow-up	Late outcomes
Wengerter et al ⁴⁰	0	1 (BTK)	0	Stroke (n = 1)	2	NR	NR
Chowdhury et al ⁴¹	0	0	0	0	1	NR	NR
Liu et al ⁴²	1	1 (TF)	0	0	0	NR	NR
Total (n)	61	34 (1 study NR)	4 (5 studies NR)	DVT/PE (n = 7) Rethrombosis (n = 2) Stroke (n = 1) AKI/hemodialysis (n = 4) Encephalopathy (n = 1) Shock (n = 2) Other limb ischemia (n = 2) Rhabdomyolysis (n = 1) HIT (n = 1) Other (n = 3) (8 studies NR)	56 (6 studies NR)	0-2 months (28 studies NR)	No new event (n = 2) Digit gangrene (n = 2) Numbness/nerve impairment (n = 2) (28 studies NR)

ARDS, Acute respiratory distress syndrome; AKI, acute kidney injury; BTK, below the knee; DVT, deep vein thrombosis; GI, gastrointestinal; HIT, heparin-induced thrombopenia; NR, not reported; PE, pulmonary embolism; TF, transfemoral.

thrombosis.³⁰ This point probably explains the high number of patients treated conservatively as a first-line treatment in this review as well. However, we found that medical treatment showed a much higher mortality risk compared with intervention, although the amputation risk was similar. In another systematic review by Putko et al,⁷⁷ a similar mortality rate was found, although the number of included cases and studies was much lower and no further meta-analysis was conducted. However, Tang et al⁷⁸ have shown that anticoagulant treatment mainly with low-molecular-weight heparin is associated with a lower mortality risk in patients with COVID-19 who have an increased coagulopathy score or high D-dimer levels. Therefore, coverage with anticoagulants is imperative in high-risk patients.

If an intervention is needed, several methods have been used for treating ALI, including thrombectomy, thrombolysis, thrombosuction, and others, yielding comparable results regarding limb salvage.⁷⁹ The choice of surgical intervention is influenced by both the clinical status of the patient and the etiology of the ALI. Data from non-COVID cases show that thrombotic rather than embolic events lead to worse outcomes.⁸⁰ In our review, we found an almost 10.5% reoperation rate and a 23.5% amputation rate, with the majority being major amputations. Other authors have also found that successful revascularization is disappointingly low in patients with COVID-19 when compared with previously reported series.⁸¹ Data indicate that continuation of anticoagulants at admission in patients already receiving such agents for other causes did not affect outcomes, even in patients undergoing operative procedures.¹² This finding underlines the strong hypercoagulant and inflammatory storm that this infection releases.

There are certain limitations to this review. First, the total number of patients included is low, and most of studies consist of case reports or case series. Second, many studies do not provide data considering the precise medical treatment or laboratory profile of patients to extract adequately powered pooled data. There was also a lack of specific a definition for the majority of comorbidities in the included studies and, therefore, these events were reported in this review as reported in the studies. Additionally, no follow-up is provided by the majority of studies. This factor is mainly due to the fact that all of studies have been published within the last year. Furthermore, reoperation rates are reported without any detail on the type of procedure in the majority of studies. Finally, data were too limited to conduct any multiregression analysis.

CONCLUSIONS

SARS-CoV-2 infection is associated with a high thrombotic risk probably by promoting a systematic inflammatory response and a hypercoagulable state. COVID-associated ALI usually presents in patients with low number of comorbidities, and it is associated with a high mortality and amputation risk. Mortality risk seems to be greater with conservative treatment compared with any intervention, although the amputation risk is similar. Future studies should focus on identifying optimal medical treatment for these patients as well as potential prognostic factors for mortality and amputation risks.

AUTHOR CONTRIBUTIONS

Conception and design: GG, AS, KF
Analysis and interpretation: GG, KS, FS

Data collection: GG, MF, KV, DV, CT, VM

Writing the article: GG, AS, MF, KV, DV, CT, VM

Critical revision of the article: GG, FS, KF

Final approval of the article: GG, AS, MF, KV, DV, CT, VM,
FS, KF

Statistical analysis: GG

Obtained funding: Not applicable

Overall responsibility: KF

REFERENCES

- Liu YC, Kuo RL, Shih SR. COVID-19: the first documented coronavirus pandemic in history. *Biomed J* 2020;43:328-33.
- Li T, Lu H, Zhang W. Clinical observation and management of COVID-19 patients. *Emerg Microbes Infect* 2020;9:687-90.
- Ejaz H, Alsrhani A, Zafar A, Javed H, Junaid K, Abdalla AE, et al. COVID-19 and comorbidities: deleterious impact on infected patients. *J Infect Public Health* 2020;13:1833-9.
- Castro RA, Frishman WH. Thrombotic complications of COVID-19 infection: a review. *Cardiol Rev* 2021;29:43-7.
- Keshavarz P, Rafiee F, Kavandi H, Goudarzi S, Heidari F, Gholamrezaezhad A. Ischemic gastrointestinal complications of COVID-19: a systematic review on imaging presentation. *Clin Imaging* 2020;73:86-95.
- Nannoni S, de Groot R, Bell S, Markus HS. Stroke in COVID-19: a systematic review and meta-analysis. *Int J Stroke* 2020;11. 1747493020972922.
- Liberati A, Altman DG, Tetzlaff J, Mulrow C, Gøtzsche PC, Ioannidis JP, et al. The PRISMA statement for reporting systematic reviews and meta-analyses of studies that evaluate healthcare interventions: explanation and elaboration. *BMJ* 2009;339:b2700.
- Stang A. Critical evaluation of the Newcastle-Ottawa scale for the assessment of the quality of nonrandomized studies in meta-analyses. *Eur J Epidemiol* 2010;25:603-5.
- Veerarasi S, Kulkarni SR, Wilson WR, Paravastu SCV. Bilateral acute lower limb ischemia secondary to COVID-19. *Vasc Endovascular Surg* 2021;55:196-9.
- Kaur P, Posimreddy S, Singh B, Qaqa F, Habib HA, Maroules M, et al. COVID-19 Presenting as acute limb ischaemia. *Eur J Case Rep Intern Med* 2020;7:001724.
- Brugliera L, Spina A, Castellazzi P, Cimino P, Arcuri P, Deriu MG, et al. Rehabilitative of COVID-19 patients with acute lower extremity Ischemia and amputation. *J Rehabil Med* 2020;52:jrm00094.
- Hanif M, Ali MJ, Haider MA, Naz S, Ahmad Z. Acute upper limb ischemia due to arterial thrombosis in a mild COVID-19 patient: a case report. *Cureus* 2020;12:e10349.
- Hasan SA, Haque A, Nazir F. Acute limb ischemia: a rare complication of COVID-19. *Cureus* 2020;12:e11488.
- Gubitosa JC, Xu P, Ahmed A, Pergament K. COVID-19-associated acute limb ischemia in a patient on therapeutic anticoagulation. *Cureus* 2020;12:e10655.
- Anwar S, Acharya S, Shabih S, Khabut A. Acute limb ischemia in COVID-19 disease: a mysterious coagulopathy. *Cureus* 2020;12:e9167.
- Singh B, Kaur P, Ajdir N, Gupta S, Maroules M. Covid-19 presenting as acute limb ischemia. *Cureus* 2020;12:e9344.
- Wang JS, Pasiaka HB, Petronic-Rosic V, Sharif-Askary B, Evans KK. Digital gangrene as a sign of catastrophic coronavirus disease 2019-related microangiopathy. *Plast Reconstr Surg Glob Open* 2020;8:e3025.
- Goldman IA, Ye K, Scheinfeld MH. Lower-extremity arterial thrombosis associated with COVID-19 is characterized by greater thrombus burden and increased rate of amputation and death. *Radiology* 2020;297:263-9.
- Sánchez JB, Cuipal Alcalde JD, Ramos Isidro R, Luna CZ, Cubas WS, Coaguila Charres A, et al. Acute limb ischemia in a Peruvian cohort infected by COVID-19. *Ann Vasc Surg* 2020;72:196-2004.
- Galanis N, Stavrakia C, Agathangelidis F, Petsatodis E, Giankoulou C, Givissis P. Coagulopathy in COVID-19 infection: a case of acute upper limb ischemia. *J Surg Case Rep* 2020;2020:rjaa204.
- Bozzani A, Arici V, Tavazzi G, Franciscone MM, Danesino V, Rota M, et al. Acute arterial and deep venous thromboembolism in COVID-19 patients: risk factors and personalized therapy. *Surgery* 2020;168:987-92.
- Mietto C, Salice V, Ferraris M, Zuccon G, Valdambri F, Piazzalunga G, et al. Acute lower limb ischemia as clinical presentation of COVID-19 infection. *Ann Vasc Surg* 2020;69:80-4.
- Baccellieri D, Bilman V, Apruzzi L, Monaco F, D'Angelo A, Loschi D, et al. A case of Covid-19 patient with acute limb ischemia and heparin resistance. *Ann Vasc Surg* 2020;68:88-92.
- Shao T, In-Bok Lee C, Jabori S, Rey J, Duran ER, Kang N. Acute upper limb ischemia as the first manifestation in a patient with COVID-19. *J Vasc Surg Cases Innov Tech* 2020;6:674-7.
- Etkin Y, Conway AM, Silpe J, Qato K, Carroccio A, Manvar-Singh P, et al. Acute arterial thromboembolism in patients with COVID-19 in the New York City area. *Ann Vasc Surg* 2021;70:290-4.
- Muhammad K, Tantawy TG, Makar RR, Olojugba O. Successful catheter-directed thrombolysis for acute lower limb ischemia secondary to COVID-19 infection. *Ann Vasc Surg* 2020;71:103-11.
- Heald M, Fish J, Lurie F. Skin manifestations of COVID-19 resembling acute limb ischemia. *J Vasc Surg Cases Innov Tech* 2020;6:514-5.
- Bellosta R, Luzzani L, Natalini C, Pegorer MA, Attisani L, Cossu LG, et al. Acute limb ischemia in patients with COVID-19 pneumonia. *J Vasc Surg* 2020;72:1864-72.
- Kaur P, Qaqa F, Ramahi A, Shamoony Y, Singhal M, Shamoony F, et al. Acute upper limb ischemia in a patient with COVID-19. *Hematol Oncol Stem Cell Ther* 2020.
- Kahlberg A, Mascia D, Bellosta R, Attisani L, Pegorer M, Socrate AM, et al. Vascular surgery during COVID-19 emergency in hub hospitals of Lombardy: experience on 305 Patients. *Eur J Vasc Endovasc Surg* 2020;61:306-15.
- Schultz K, Wolf JM. Digital ischemia in COVID-19 patients: case report. *J Hand Surg Am* 2020;45:518-22.
- Vacirca A, Faggioli C, Pini R, Teutonico P, Pilato A, Gargiulo M. Unheralded lower limb threatening ischemia in a COVID-19 patient. *Int J Infect Dis* 2020;96:590-2.
- Fan BE, Chia YW, Sum CLL, Kuperan P, Chan SSW, Ling LM, et al. Global haemostatic tests in rapid diagnosis and management of COVID-19 associated coagulopathy in acute limb ischaemia. *J Thromb Thrombolysis* 2020;50:292-7.
- Perini P, Nabulsi B, Massoni CB, Azzarone M, Freyrie A. Acute limb ischaemia in two young, non-atherosclerotic patients with COVID-19. *Lancet* 2020;395:1546.
- Kashi M, Jacquin A, Dakhil B, Zaimi R, Mahé E, Tella E, et al. Severe arterial thrombosis associated with Covid-19 infection. *Thromb Res* 2020;192:75-7.
- Baeza C, González A, Torres P, Pizzamiglio M, Arribas A, Aparicio C. Acute aortic thrombosis in COVID-19. *J Vasc Surg Cases Innov Tech* 2020;6:483-6.
- Garg K, Barfield ME, Pezold ML, Sadek M, Cayne NS, Lugo J, et al. Arterial thromboembolism associated with COVID-19 and elevated D-dimer levels. *J Vasc Surg Cases Innov Tech* 2020;6:348-51.
- Thompson O, Pierce D, Whang D, O'Malley M, Geise B, Malhotra U. Acute limb ischemia as sole initial manifestation of SARS-CoV-2 infection. *J Vasc Surg Cases Innov Tech* 2020;6:511-3.
- Levolger S, Bokkers RPH, Wille J, Kropman RHJ, de Vries JPM. Arterial thrombotic complications in COVID-19 patients. *J Vasc Surg Cases Innov Tech* 2020;6:454-9.
- Wengerter SP, Wengerter KR, Masoudpoor H, Sagarwala A, Karim O, Rao N, et al. Acute aortoiliac and infrainguinal arterial thrombotic events in four patients diagnosed with the novel coronavirus 2019. *J Vasc Surg Cases Innov Tech* 2020;6:698-702.
- Chowdhury YS, Mitre CA, Rotella VE, Garg K, Lee DK, Belligund P, et al. Extensive peripheral arterial thrombosis in a patient with SARS-CoV-2 infection. *Am J Med Case Rep* 2020;8:486-90.
- Liu Y, Chen P, Mutar M, Hung M, Shao Z, Han Y, et al. Ischemic necrosis of lower extremity in COVID-19: a case report. *J Atheroscler Thromb* 2021;28:90-5.
- Baril DT, Ghosh K, Rosen AB. Trends in the incidence, treatment, and outcomes of acute lower extremity ischemia in the United States Medicare population. *J Vasc Surg* 2014;60:669-77.
- Levi M, Thachil J, Iba T, Levy JH. Coagulation abnormalities and thrombosis in patients with COVID-19. *Lancet Haematol* 2020;7: 438-40.
- Klok FA, Kruij MJHA, van der Meer NJM, Arbous MS, Gommers DAMPJ, Kant KM, et al. Incidence of thrombotic complications in critically ill ICU patients with COVID-19. *Thromb Res* 2020;191:145-7.

46. Shah A, Donovan K, McHugh A, Pandey M, Aaron L, Bradbury CA, et al. Thrombotic and haemorrhagic complications in critically ill patients with COVID-19: a multicentre observational study. *Crit Care* 2020;24:561.
47. Eliason JL, Wainess RM, Proctor MC, Dimick JB, Cowan JA Jr, Upchurch GR Jr, et al. A national and single institutional experience in the contemporary treatment of acute lower extremity ischemia. *Ann Surg* 2003;238:382-9.
48. Hemingway J, Emanuels D, Aarabi S, Quiroga E, Tran N, Starnes B, et al. Safety of transfer, type of procedure, and factors predictive of limb salvage in a modern series of acute limb ischemia. *J Vasc Surg* 2019;69:1174-9.
49. Lodigiani C, Iapichino G, Carenzo L, Cecconi M, Ferrazzi P, Sebastian T, et al. Venous and arterial thromboembolic complications in COVID-19 patients admitted to an academic hospital in Milan, Italy. *Thromb Res* 2020;191:9-14.
50. Bissacco D, Grassi V, Lomazzi C, Domanin M, Bellosta R, Piffaretti G, et al. Is there a vascular side of the story? Vascular consequences during COVID-19 outbreak in Lombardy, Italy. *J Card Surg* 2020;36:1677-82.
51. de Roquetaillade C, Chousterman BG, Tomasoni D, Zeitouni M, Houdart E, Guedon A, et al. Unusual arterial thrombotic events in Covid-19 patients. *Int J Cardiol* 2021;323:281-4.
52. Albitar O, Ballouze R, Ooi JP, Sheikh Ghadzi SM. Risk factors for mortality among COVID-19 patients. *Diabetes Res Clin Pract* 2020;166:108293.
53. McGonagle D, Sharif K, O'Regan A, Bridgwood C. The role of cytokines including interleukin-6 in COVID-19 induced pneumonia and macrophage activation syndrome-like disease. *Autoimmun Rev* 2020;19:10253.
54. Zeng F, Huang Y, Guo Y, Yin M, Chen X, Xiao L, et al. Association of inflammatory markers with the severity of COVID-19: a meta-analysis. *Int J Infect Dis* 2020;96:467-74.
55. Liu B, Li M, Zhou Z, Guan X, Xiang Y. Can we use interleukin-6 (IL-6) blockade for coronavirus disease 2019 (COVID-19)-induced cytokine release syndrome (CRS)? *J Autoimmun* 2020;111:102452.
56. Mehta P, McAuley DF, Brown M, Sanchez E, Tattersall RS, Manson JJ. COVID-19: consider cytokine storm syndromes and immunosuppression. *Lancet* 2020;395:1033-4.
57. Terpos E, Ntanasis-Stathopoulos I, Elalamy I, Kastritis E, Sergentanis TN, Politou M, et al. Hematological findings and complications of COVID-19. *Am J Hematol* 2020;95:834-47.
58. Tang N, Li D, Wang X, Sun Z. Abnormal coagulation parameters are associated with poor prognosis in patients with novel coronavirus pneumonia. *J Thromb Haemost* 2020;18:844-7.
59. Magro C, Mulvey JJ, Berlin D, Nuovo G, Salvatore S, Harp J, et al. Complement associated microvascular injury and thrombosis in the pathogenesis of severe COVID-19 infection: a report of five cases. *Transl Res* 2020;220:1-13.
60. Wichmann D. Autopsy findings and venous thromboembolism in patients with COVID-19. *Ann Intern Med* 2020;173:1030.
61. Alonso MN, Mata-Forte T, García-León N, Vullo PA, Ramirez-Olivencia G, Estébanez M, et al. Incidence, characteristics, laboratory findings and outcomes in acro-ischemia in COVID-19 patients. *Vasc Health Risk Manag* 2020;16:467-78.
62. Ackermann M, Verleden SE, Kuehnel M, Haverich A, Welte T, Laenger F, et al. Pulmonary vascular endothelialitis, thrombosis, and angiogenesis in Covid-19. *N Engl J Med* 2020;383:120-8.
63. Varga Z, Flammer AJ, Steiger P, Haberecker M, Andermatt R, Zinkernagel AS, et al. Endothelial cell infection and endotheliitis in COVID-19. *Lancet* 2020;395:1417-8.
64. Goeijenbier M, van Wissen M, van de Weg C, Jong E, Gerdes VE, Meijers JC, et al. Review: viral infections and mechanisms of thrombosis and bleeding. *J Med Virol* 2012;84:1680-96.
65. Zhang Y, Xiao M, Zhang S, Xia P, Cao W, Jiang W, et al. Coagulopathy and antiphospholipid antibodies in patients with Covid-19. *N Engl J Med* 2020;382:38.
66. Galli L, Gerdes VE, Guasti L, Squizzato A. Thrombosis associated with viral hepatitis. *J Clin Transl Hepatol* 2014;2:234-9.
67. Jackson BS, Pretorius E. Pathological clotting and deep vein thrombosis in patients with HIV. *Semin Thromb Hemost* 2019;45:132-40.
68. Zhang L, Yan X, Fan Q, Liu H, Liu X, Liu Z, et al. D-dimer levels on admission to predict in-hospital mortality in patients with Covid-19. *J Thromb Haemost* 2020;18:1324-9.
69. Yin S, Huang M, Li D, Tang N. Difference of coagulation features between severe pneumonia induced by SARS-CoV2 and non-SARS-CoV2. *J Thromb Thrombolysis* 2020;3:1-4.
70. Xiong M, Liang X, Wei YD. Changes in blood coagulation in patients with severe coronavirus disease 2019 (COVID-19): a meta-analysis. *Br J Haematol* 2020;189:1050-2.
71. Lippi G, Favaloro EJ. D-dimer is associated with severity of coronavirus disease 2019: a pooled analysis. *Thromb Haemost* 2020;120:876-8.
72. Al-Ani F, Chehade S, Lazo-Langner A. Thrombosis risk associated with COVID-19 infection. A scoping review. *Thromb Res* 2020;192:152-60.
73. Bikdeli B, Madhavan M, Jimenez D, Chuich T, Dreyfus I, Driggin E, et al. COVID-19 and thrombotic or thromboembolic disease: implications for prevention, antithrombotic therapy, and follow-up: JACC state-of-the-art review. *J Am Coll Cardiol* 2020;75:2950-73.
74. Avila J, Long B, Holladay D, Gottlieb M. Thrombotic complications of COVID-19. *Am J Emerg Med* 2021;39:213-8.
75. Artifoni M, Danic G, Gautier C, Gicquel P, Boutoille D, Raffi F, et al. Systematic assessment of venous thromboembolism in COVID-19 patients receiving thromboprophylaxis: incidence and role of D-dimer as predictive factors. *J Thromb Thrombolysis* 2020;50:211-6.
76. Han H, Yang L, Liu R, Liu F, Wu KL, Li J, et al. Prominent changes in blood coagulation of patients with SARS-CoV-2 infection. *Clin Chem Lab Med* 2020;58:1116-20.
77. Putko RM, Bedrin MD, Clark DM, Piscocoy AS, Dunn JC, Nesti LJ. SARS-CoV-2 and limb ischemia: a systematic review. *J Clin Orthop Trauma* 2021;12:194-9.
78. Tang N, Bai H, Chen X, Gong J, Li D, Sun Z. Anticoagulant treatment is associated with decreased mortality in severe coronavirus disease 2019 patients with coagulopathy. *J Thromb Haemost* 2020;18:1094-9.
79. Veenstra EB, van der Laan MJ, Zeebregts CJ, de Heide EJ, Kater M, Bokkers RPH. A systematic review and meta-analysis of endovascular and surgical revascularization techniques in acute limb ischemia. *J Vasc Surg* 2020;71:654-68.e3.
80. Torrealba JI, Osman M, Kelso R. Hypercoagulability predicts worse outcomes in young patients undergoing lower extremity revascularization. *J Vasc Surg* 2019;70:175-80.
81. Piffaretti C, Angrisano A, Franchin M, Ferrario M, Rivolta N, Bacuzzi A, et al. Risk factors analysis of thromboembolism for acute thromboembolic lower extremity ischemia in native arteries. *J Cardiovasc Surg* 2018;59:810-6.

Submitted Mar 30, 2021; accepted Jul 27, 2021.