distance between people (n = 7); three provided hand sanitizers, only one provided gloves. Regarding employees, in four of the eight stores observed the cashiers wore a visor or mask/glasses and gloves; in seven stores there was an acrylic separator wall between them and the customers. Store operators wore gloves in three out of eight outlets, but in only two cases they used a visor or a mask. Of the 78 customers observed, 47 (60.3%) were women and the majority were between 18 and 65 years old (n = 60; 76.9%). In the total sample, 26.9% (n = 21) used a surgical mask and 0.05% (n = 4) used a handmade mask or scarf. No customer wore a visor. Gloves were the most commonly used equipment by customers (n = 25; 32.1%).

To sum up, compliance with prevention measures recommended to avoid COVID-19 contamination in retail food stores in Braga varied widely between outlets in April 2020. There were examples of excellent practice, but also of poor practice. Only one store met all recommended standards. The situation is more worrying in mini markets, where none of the recommendations were being followed. The main shortcomings, in general, were the absence of protective equipment for the employees, and lack of hand sanitizer for customers. According to recent information from the European Centre for Disease Prevention and Control, one of the best preventive practices to reduce the transmission of COVID-19 is the generalized use of face masks as a complementary measure to safety distance, proper education being fundamental for their safe use.

Ethical disclosures

Protection of human and animal subjects
The authors declare that no experiments were performed on humans or animals for this study.

Confidentiality of data
The authors declare that the procedures followed were in accordance with the regulations established by the Commission for Clinical and Ethical Research and in accordance with the Helsinki Declaration of the World Medical Association.

Right to privacy and informed consent
The authors have obtained the written informed consent of the patients or subjects mentioned in the article. The corresponding author is in possession of this document.

Conflict of interest
The authors have no conflicts of interest to declare.

Authors’ contributions
Precioso conceived this study, collected the data, designed, and carried out statistical analysis. Samorinha and Precioso wrote the first draft of the manuscript. Samorinha carried out statistical analysis and revised the manuscript critically. All authors contributed substantially to the interpretation of data, critical discussion, and revision of the manuscript, and approved its final version.

References
1. World Health Organization. https://www.who.int/emergencies/diseases/novel-coronavirus-2019/technical-guidance/naming-the-coronavirus-disease-(covid-19)-and-the-virus-that-causes-it, 2020 [accessed 12 June 2020].
2. Serviço Nacional de Saúde. https://www.sns24.gov.pt/tema/doencas-infecciosas/covid-19/#sec-0, 2020 [accessed 12 June 2020].
3. Direcção-Geral da Saúde. https://www.dgs.pt/diretrizes-da-dgs/orientacoes-e-circulares-informativas/orientacao-n-0142020-de-21032020-pdf.aspx, 2020 [accessed 12 June 2020].
4. Centres for Disease Control and Prevention. https://www.cdc.gov/coronavirus/2019-ncov/prevent-getting-sick/diy-cloth-face-coverings.html, 2020 [accessed 12 June 2020].

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Pulmonary artery thrombosis in COVID-19 patients

Dear Editor,

Arising in China in the winter of 2019, COVID-19 (caused by the SARS-CoV-2 virus) has caused a global pandemic and severely stressed medical systems across the world.

Although knowledge about this novel coronavirus is still emerging, the most common reason for hospitalization of COVID-19 patients is severe respiratory distress.1 COVID-19 has been accurately described as the cause for a proinflammatory and hypercoagulable state with marked elevations seen in Lactate Dehydrogenase, Ferritin, C-reactive protein, D-Dimer, and Interleukin levels.2

The inflammatory response, including production of inflammatory cells and cytokines, induces a procoagulant effect and diffuse endothelial damage that predisposes thrombotic vascular lesions and Disseminated Intravascular Coagulation (DIC).3 D-Dimer is related to the severity of the disease and an increased value is associated with the worst prognosis. Retrospective studies demonstrated that patients admitted to
Intensive Care Unit (ICU) had an elevated D Dimer value and, in this setting, some Authors recommended a therapeutic heparin doses for the patients with higher values.\(^4\)

A recent ICU observation reported an increased risk of Pulmonary Embolism (PE) in COVID-19 compared to the historical control group even in patients that had undergone the Low Molecular Weight Heparin (LMWH) prophylaxis.\(^5\)

We evaluated 138 patients with COVID-19 admitted to our Institution between March 2020 and May 2020. All patients were COVID-19 positive according to clinical diagnostic criteria reverse-transcription–polymerase chain-reaction (RT-PCR) and Chest Thoracic tomography. On admission, most of them were haemodynamically stable (78%) and febrile (87%). During hospitalization, some developed progressive respiratory failure and received oxygen supplementation (41%). Four of them were started on Continuous Positive Airways Pressure (CPAP) but two died because of worsening Respiratory Failure.

All patients were treated with hydroxychloroquine (400 mg/day), darunavir/ritonavir (800/100 mg/day) and enoxaparin (4000 UI/day). Some patients (26 pts) received additional therapy with IL-6 and IL-1 antagonist (20%). Every three days after their hospitalization, laboratory exams with inflammatory and coagulation parameters (INR, activated partial thromboplastin time, platelet count, fibrinogen, D-Dimer) were repeated.

In patients with progressive elevation of D-Dimer of over three times the normal value (from 1822 to 5911 μg/mL), we performed a Computed Tomography Pulmonary Angiography (CTPA) and a Doppler Ultrasound (DU) of the lower limbs. The tests were done during the second week of their hospitalization (12,3 ± 3,2 days).

We identified eleven patients with high D-Dimer value, nine of them (6,7%) had signs of Pulmonary Artery Thrombosis (PAT) without Deep Venous Thrombosis (DVT). The lesions were distributed bilaterally at the lower arterial branches (Figure 1). None of the nine patients experienced an objective respiratory worsening and those in oxygen therapy (5 patients) maintained a constant flow. Therapeutic anticoagulation was started with subcutaneous enoxaparin (1 mg/kg) twice daily, followed by warfarin. All patients were discharged home: length of hospital stay (LOS) 21,1 ± 3,7 days.

In Table 1 the laboratory parameters were compared between patients with and without PAT: five days after admission, only D-Dimer value and lymphocyte count were significantly different between the two groups.

Parameters are expressed as mean ± SD and statistically evaluated by Student T test. \(P < 0,05\) was considered statistically significant.

Coagulopathic disorders are significantly increased in COVID-19 patients, especially among those with severe dis-

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**Table 1** Laboratory parameters in patients with Pulmonary Artery Thrombosis (PAT) and the patients without PAT.

|                | All Pts admission | PAT pts admission | All pts 5 days | PAT pts 5 days | All Pts 10 days | PAT pts 10 days |
|----------------|-------------------|-------------------|---------------|---------------|----------------|---------------|
| C-Reactive Protein (mg/l) | 54,6 ± 58,1 | 54,6 ± 58,1 | 44,9 ± 65,1 | 54,6 ± 58,1 | 27,0 ± 25,5 | 49,5 ± 27,6 |
| Lymphocite (mmc) | 1294 ± 761,3 | 1260 ± 414,5 | 1465 ± 657,6 | 1075 ± 219,2* | 1762 ± 858,6 | 1290 ± 336,7* |
| Platelets (mmc) | 250,000 ± 84,000 | 263,000 ± 147,000 | 300,000 ± 114,000 | 252,800 ± 52,900 | 352,300 ± 102,900 | 289,000 ± 113,000 |
| Fibrinogen (mg/dl) | 432,3 ± 114,1 | 530 ± 121,6 | 428,5 ± 138,2 | 426,0 ± 14,1 | 379,6 ± 100,4 | 466,1 ± 142,1 |
| LDH (U/l) | 264,4 ± 101,3 | 326,2 ± 121,9 | 295,2 ± 120,6 | 314,5 ± 37,5 | 314,5 ± 37,5 | 327,5* |
| INR | 1 ± 0,1 | 1 ± 0,01 | 1,2 ± 0,15 | 1,05 ± 0,1 | 1,2 ± 0,1 | 1,1 ± 0,1 |
| D-Dimer Ug/l | 465,1 ± 121,7 | 627,4 ± 178,6 | 538,3 ± 102,8 | 2143,6 ± 327,5* | 576,9 ± 169,1 | 1764,9 ± 227,5* |

* \(P < 0,05\).
ease. Several mechanisms combine systemic inflammation with alterations of coagulation in COVID-19 patients. In severe or critically ill patients, the endothelial cells are damaged and release a large amount of inflammatory mediators that may predispose vascular thrombosis. A study performed in ICU setting reported an increased risk of Pulmonary Embolism (PE) in COVID-19 patients treated with Low Molecular Weight Heparin (LMWH) prophylaxis compared to the historical COVID 19 negative control group. Another Study reported that the prevalence of Venous Thromboembolism Events (VTE) was higher in ICU compared to general wards patients: 47% vs 3%. High blood values of the procoagulant factor levels including fibrinogen and D-dimers have been associated with the worst prognosis and higher mortality.

Kaminetzky and coworkers compared the results of a cohort of 62 patients who underwent CTPA for suspected PE prior to the first case of COVID 19, with 62 patients COVID 19 positive. CTPA was positive for PE in 37% of COVID 19 patients (14.5% in pre COVID patients), D-Dimer was associated with a higher prevalence of thromboembolic events and correlated with the degree of PE severity.

In a group of patients admitted to non-ICU wards, DU failed to detect DVT independently of the severity of their condition and length of in-hospital bed rest. The Authors observed that this is apparently in contrast with the relatively frequent reports of PE in hospitalized COVID-19 patients It is possible that local thrombi in the lungs may be the cause of pulmonary arterial manifestations. In this paper we reported COVID-19 patients with interstitial pneumonia admitted in our non-ICU Ward.

During the course of hospitalization, in eleven of them, we observed a progressive increase of D-Dimer over three times the normal value, associated with low or normal values of other coagulation or inflammatory blood parameters (CRP, LDH, Ferritin, fibrinogen, INR, aPTT).

Nine CTPA demonstrated a distal thrombosis of the lower pulmonary arterial branches. The mainly basal localization where the pulmonary inflammation is most diffuse and the loss of signs of DVT may suggest a pulmonary thrombosis rather than an embolism.

It is noteworthy that no patients had any signs of respiratory worsening and some of them did not receive oxygen therapy and they were breathing room air.

In conclusion, we described patients with moderate disease who developed a pulmonary vascular injury strictly correlated with an elevation of D-Dimer values. This parameter may help clinicians in identifying COVID-19 stable patients at risk of concurrent pulmonary artery thrombosis.

Further studies will need to better define the meaning of these preliminary observations.

Conflicts of interest

The authors have no conflicts of interest to declare.

References

1. Wang D, Hu B, Hu C, Fangfang Z, Xing L, Jing Z, et al. Clinical characteristics of 138 hospitalized patients with 2019 novel coronavirus–infected pneumonia in Wuhan, China. JAMA. 2020;323(11):1061-9, http://dx.doi.org/10.1001/jama.2020.1583.
2. Connors JM, Levy JH. Thromboinflammation and the hypercoagulability of COVID-19. J Thromb Haemost. 2020;18(7):1559–61, http://dx.doi.org/10.1111/jth.14849.
3. Terpos E, Ntanasis-Stathopoulos I, Elalamy I, Kastritis E, Ser gentanis TN, Politou M, et al. Ematological findings and complications of COVID-19. Am J Emalol.2020;95(7):834-47, http://dx.doi.org/10.1002/aem.25829.
4. 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(5):1094–9, http://dx.doi.org/10.1111/jth.14817.
5. Xu Z, Shi L, Wang Y, Zhang J, Huang L, Zhang C, et al. Pathological findings of COVID-19 associated with acute respiratory distress syndrome. Lancet Resp Med. 2020;8(4):420–2, http://dx.doi.org/10.1016/S2213-2600(20)30076-X.
6. Lillicrap D. Disseminated intravascular coagulation in patients with 2019–nCoV pneumonia. J Thromb Haemost. 2020;18(Apr (4)):786–7, http://dx.doi.org/10.1111/jth.14781.
7. Poissy J, Goutay J, Morgan Caplan M, Parmentier E, Duburoc T, Lassalle F, et al. Pulmonary embolism in COVID-19 patients: awareness of an increased prevalence. Circulation. 2020;142:184–6, http://dx.doi.org/10.1161/CIRCULATIONAHA.120.047430.
8. Middeldorp S, Coppens M, Van Haaps TF, Foppen M, Vlaar AP, Muller M, et al. Incidence of venous thromboembolism in hospitalized patients with COVID-19. J Thromb Haemost. 2020, http://dx.doi.org/10.1111/jth.14888.
9. Kaminetzky M, Moore W, Fansiwala K, Babb JS, Kaminetzky D, Horowitz L, et al. Pulmonary embolism on CTPA in COVID-19 patients. Cardiothoracic Imaging. 2020;2(4):220–9, http://dx.doi.org/10.1148/ryct.202000308.
10. Klok FA, Kruip MJHA, van der Meer NJM, Arbous MS, Gommers PJ, Kant KH, et al. Incidence of thrombotic complications in critically ill ICU patients with COVID-19. Thromb Res. 2020;191:145–7, http://dx.doi.org/10.1016/j.thromres.2020.04.013.
11. Cattaneo M, Bertinato EM, Birocchi S, Brizio C, Malavolta D, Manzoni M, et al. Pulmonary embolism or pulmonary thrombosis in COVID-19? Is the recommendation to use high-dose heparin for thromboprophylaxis justified? Thromb Haemost. 2020, http://dx.doi.org/10.1055/s-0040-1712097 [in press].

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