Antithrombotic drugs with adjuvant action against COVID-19

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ABSTRACT

Waiting for the vaccine and/or the best antiviral treatment for coronavirus infection disease 2019 (COVID-19), after its outbreak in China at the beginning of 2020 and its viral diffusion around the world in the following weeks, several drugs have been suggested for their potential adjuvant support against infection. Several drugs have been suggested to have a potential ancillary antiviral role. Circulating proteins, in particular proteases and peptidases regulated by drug functions, may interact with well-known drugs like anticoagulants, antihypertensives, antiserotoninergics, and so also with viral proteases. We here report a brief list of these drugs (i.e., heparinoids, flavonoids, antiplatelets, anticoagulants) that may interact with COVID-19.

Introduction

Venous thromboembolism (VTE) has been frequently reported as a clinical complication of hospitalization of patients affected by coronavirus infection disease 2019 (COVID-19) in particular in Intensive Care Unit.1,2 Moreover, after the viral diffusion of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) from China to other countries in the world, several reports testified that alteration of hemostasis as the increase of fibrinogen and D-dimer might be associated with poor outcome not only of COVID-19 per se but also of the associated VTE.3-5

For this reason, the benefit of antithrombotic drugs during COVID-19 was discovered.6 Antithrombotic drugs, in fact, may have specific antiviral and anti-inflammatory actions as testified in vitro besides their attested antithrombotic actions.

So, antithrombtics may interact with infections and with their most common complications (i.e. VTE). Heparinoids, may exert their actions because of their ability to interact with viral proteases7 and because of their antithrombotic actions.6 However, other anticoagulants can exert their action, only preventing thrombotic complications as well as aspirin and other antithrombotic drugs.

So in this report, we summarized the action of several drugs able to improve the outcome of COVID-19 infection in different ways.

Flavonoids

Flavonoids are natural products; they have a polyphenolic structure widely found in fruits and vegetables.8 They have various biochemical and antioxidant effects that may be useful during various diseases such as cancer, Alzheimer’s disease, and atherosclerosis.9,10

Intriguingly, some flavonoids also have antiviral activity, as demonstrated in vitro. Their main antiviral action is related to the ability to inhibit viral proteases, in particular SARS-CoV 3-C-Like-protease.9,10

Furthermore, they may exert their antiviral action because of the affinity with oligosaccharides of the external wall of several cells that are responsible for the interaction between virus and host, in particular, such fucosylated oligosaccharides.
Heparan sulfate

Heparan sulfate (HS) is an abundant cell-surface glycosaminoglycan (GAG) that is also present in the extracellular matrix and can bind several viruses; however, other GAGs present on cellular surfaces as chondroitin sulfate showed similar properties.

From a pathophysiological point of view, HS is also abundant in the respiratory tract, and it plays a role as a binding factor for coronaviruses with tropism to bronchitis and other respiratory infections.

These actions may explain, in part, the extended tropism of this virus for cells of the respiratory tract.

Moreover, HS is also present on the surface of endothelial cells, in particular, next to alveolar-capillary areas, so the capacity of SARS-CoV-2 to induce damages to lungs is due also to this property.

Therefore, viral actions and viral capacity of SARS-CoV-2 to bind host cells may slow down in the presence of high doses of HS, and these are really important for the pathophysiology of COVID-19 infection.

Heparins

Unfractionated heparins and low molecular weight heparins have a similar structure with HS, so an action similar to HS may be hypothesized during infection of coronaviruses.

Furthermore, heparins can exert their activity, increasing the action of several anti-proteases as antithrombin and other serpin family members.

Moreover, heparins and HS are also able to interact not only with clotting factors but also with endothelial cells, so their role in the pathophysiology of microvascular thrombosis that is described in COVID-19 infection is particularly important.

In this field, in fact, as a hypercoagulable state has been testified in several articles in patients with COVID-19 infection, pulmonary embolism and other types of thromboses may be associated with morbidity and mortality of these patients. So, heparins also assume an important role also for the thromboprophylaxis and prevention of venous thromboembolism in this infection.

For this reason, prophylactic anticoagulation with LMWH as early as possible is suggested for patients affected by COVID-19 to prevent thrombotic events and organ damage. This recommendation has been recently published in a preliminary International Society on Thrombosis and Haemostasis (ISTH). Furthermore, after several clinical round tables, also the use of intermediated doses of LMWH has been recently reported as safe and effective in a clinical study. Of course, patients with anticoagulant contraindications should not do it.

Fondaparinux

Fondaparinux is the synthetic pentasaccharide sequence with antithrombotic actions present in all heparins. In daily clinical practice, fondaparinux and enoxaparin have the same clinical indication for the prevention of VTE. Nevertheless, although the action of heparins toward SARS-CoV-2 has been found in vitro and in vivo, data concerning fondaparinux lack in the Literature. Italian Authors recently suggested a good antithrombotic action of fondaparinux in VTE prevention in COVID-19 in the FONDENOXAVID study (data not published).

Aspirin and other antiplatelets

A specific antiviral action of aspirin toward SARS-CoV-2 has been found neither in vitro nor in vivo, but the use of aspirin for patients with COVID-19 seems to have several advantages.

The majority of patients admitted to hospital for COVID-19 in the world, in fact, showed age older than 50 yy, and for this reason, they were already taking cardiovascular and antithrombotic drugs for the secondary prevention of cardiovascular diseases.

In particular, an Italian cohort of patients admitted to the hospital for COVID-19 were on treatment with antiplatelets for secondary prevention of atherothrombotic disease (i.e., acute myocardial infarction or ischaemic stroke). Interestingly, in this study, the rate of admitted patients for COVID-19 ongoing antiplatelets was higher than the one of patients taking anticoagulants. Furthermore, in this cohort of patients, aspirin was the most frequent antiplatelet drug, while dual antiplatelet therapy was present for a minority of patients. Interestingly, patients taking aspirin during hospitalization did not show increased mortality for thrombotic disorders.

So, based on the data of this Italian study, the suggestion not to withdraw the chronic use of aspirin has been made; the benefit of its action was based on the prevention of ischemic stroke and acute coronary syndrome, and also on the safety for the low rate of bleeding complications.

Furthermore, the safety of aspirin during the COVID-19 outbreak has also been reported in pregnant women affected by this viral infection.

Warfarin and direct oral anticoagulants

A specific action of oral anticoagulants toward SARS-CoV-2 is not present. However, for the frequent association of chronic arrhythmias or paroxysmal arrhythmias as atrial fibrillation, the use of anti-vitamin K (AVK) or direct oral anticoagulants (DOACs) has been found safe during the COVID-19 outbreak.
Furthermore, the use of AVK or DOACs is also associated with a reduced risk of developing VTE during COVID-19.22

Nevertheless, in this clinical setting, the administration of DOACs to these patients should be thoroughly performed because a possible drug-drug interaction between DOACs and antivirals.23

Furthermore, the use of any type of anticoagulant therapy is not associated with developing ARDS or death during hospitalization for COVID-19.22

So, the use of any oral anticoagulants (i.e., AVK or DOACs) may help escape different types of embolism during COVID-19, as venous thromboembolism due to hypercoagulable state and/or cardioembolism due to the frequent comorbidities as atrial fibrillation or other cardiovascular diseases.22

**Bleedings**

Although COVID-19 has been associated with thrombophilia and thrombosis since first reports, recently, a clinical study underlines also a relevant rate of bleedings. In this study, the incidence of thrombotic complications was similar to the rate of major bleedings.24

These data seem to be different from those previously published from other Authors.

In the Italian cohort of patients affected by COVID-19 and treated with antiplatelets, the number of bleedings was inconsistent.21 Also, a clinical registry from Austria on gastrointestinal bleedings reported a reduced rate of bleeding events during the outbreak, probably also related to home rest for lockdown.25 So, because the international scientific societies have recommended prophylactic LMWH without an excess of bleedings at standard doses, and at intermediate doses, also higher rates of bleeding events during the outbreak, probably also related to home rest for lockdown.25

Of course, bleedings may occur in such patients and could be explained by the presence of comorbidities with the trend to bleed of affected patients and also by the use of antithrombotic drugs.

So, a clinical evaluation of bleeding risk in patients affected by COVID-19 should always be considered when the administration of antithrombotic is ongoing as well as for other acute medical illnesses.

**Conclusions**

Several drugs have been suggested to improve the outcome of COVID-19 infection, particularly anticoagulants, for the associated risk to develop VTE during infection. This association is one of the reasons for which this outbreak has been more aggressive than the previous coronavirus epidemics (i.e., SARS and MERS). The target of these drugs may be viral proteases or other proteases that may interfere with the clustering system or inflammatory system. Furthermore, because cardiovascular diseases with a trend to cardioembolism have been frequently reported in all populations of infected patients, also other anticoagulants as AVK or DOACs showed their useful actions during the COVID-19 outbreak.

Furthermore, additional useful actions have been demonstrated by flavonoids and heparins because of their ability to counteract the viral action on the cellular surface and the extracellular matrix. In conclusion, waiting for specific studies that may help understand the real power of these drugs in the prevention of mortality of patients with COVID-19, we strongly encourage to use the better antithrombotic strategy to prevent any thrombotic complication during COVID-19.

**References**

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