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► **To cite this version:**

Asli Gelincik, Knut Brockow, Gülfem Çelik, Inmaculada Doña, Cristobalina Mayorga, et al.. Diagnosis and management of the drug hypersensitivity reactions in Coronavirus disease 19. *Allergy*, 2020, 10.1111/all.14439 . hal-02934360

HAL Id: hal-02934360

<https://hal.sorbonne-universite.fr/hal-02934360>











Submitted on 9 Feb 2024

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Diagnosis and management of the drug hypersensitivity reactions in Coronavirus disease 19: An EAACI Position Paper

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Abstract

Coronavirus disease 2019 (COVID-19), a respiratory tract infection caused by a novel human coronavirus, the severe acute respiratory syndrome coronavirus 2, leads to a wide spectrum of clinical manifestations ranging from asymptomatic cases to patients with mild and severe symptoms, with or without pneumonia. Given the huge influence caused by the overwhelming COVID-19 pandemic affecting over three million people worldwide, a wide spectrum of drugs is considered for the treatment in the concept of repurposing and off-label use. There is no knowledge about the diagnosis and clinical management of the drug hypersensitivity reactions that can potentially occur during the disease. This review brings together all the published information about the diagnosis and management of drug hypersensitivity reactions due to current and candidate off-label drugs and highlights relevant recommendations. Furthermore, it gathers all the dermatologic manifestations reported during the disease for guiding the clinicians to establish a better differential diagnosis of drug hypersensitivity reactions in the course of the disease.

KEYWORDS

COVID-19, desensitization, drug hypersensitivity reactions, SARS-CoV-2

1 | INTRODUCTION

Coronavirus disease 2019 (COVID-19) is a respiratory tract infection caused by a novel member of human coronavirus, the severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2).¹ It causes a wide spectrum of clinical manifestations ranging from asymptomatic cases to patients with mild, uncomplicated illness and severe cases, with or without pneumonia.² Hospitalization and oxygen support, and admission to an intensive care unit are required in 14% and 5% of the patients, respectively.¹ Gastrointestinal symptoms and positive viral nucleic acid testing on rectal swabs are considered as indicators of infection in digestive system and fecal-oral transmission of COVID-19.³ Moreover, skin symptoms, including exanthems, may appear during the evolution of the disease leading to differential diagnosis with drug hypersensitivity reactions (DHRs).⁴

In critically ill patients, COVID-19 can be complicated by acute respiratory distress syndrome (ARDS), septic shock, and multi-organ dysfunction syndrome.¹ In such patients, in response to viral infection, the excessive activation and expansion of T lymphocytes and macrophages lead to an overproduction of cytokines, which causes a cytokine storm and a hyperinflammatory state.^{5,6} Acute hyperinflammation may activate coagulation cascade and inhibit fibrinolytic reaction, thus promoting thrombosis. Coagulopathy and thrombocytopenia are serious complications which increase the risk of hemorrhage and thrombosis and progress to disseminated intravascular coagulation (DIC).⁷

The periodically updated World Health Organisation interim guidance allows reliable comparison of investigational therapeutic interventions as part of randomized controlled trials, provides recommendations for the management, and forms the basis of many institutional or national protocols.¹ Unfortunately, none of the drugs used for COVID-19 have been proven to be truly effective yet; besides, no specific antiviral drugs have been approved for COVID-19 by health authorities.^{8,9} At the moment, there is no specific treatment for COVID-19, and standard practice of care focuses on treating the clinical symptoms with supportive care.¹

In this review, diagnosis and management of DHRs, which are expected to be caused by current or candidate repurposed and off-label drugs used for COVID-19 treatment mostly based on prior knowledge, are discussed.^{8,10,11} Drugs in this review are classified into four groups according to their potential roles in different phases of the disease as antiviral drugs, antiviral and/or immunomodulatory drugs used in viral pneumonia; anti-cytokine and anti-inflammatory drugs considered during macrophage activation syndrome (MAS) and cytokine storm; anti-inflammatory drugs in ARDS; and anti-aggregant and anti-coagulant drugs in coagulopathy (Figure 1). Information of DHRs due to the use of additional drugs for various purposes can be found in the relevant European Academy of Allergy and Clinical Immunology (EAACI) resources.¹²⁻²⁰

Since emerging recent findings are dynamically changing the clinical interventions, it is expected that the list of drugs determined according to current knowledge may change with upcoming recommendations in future.

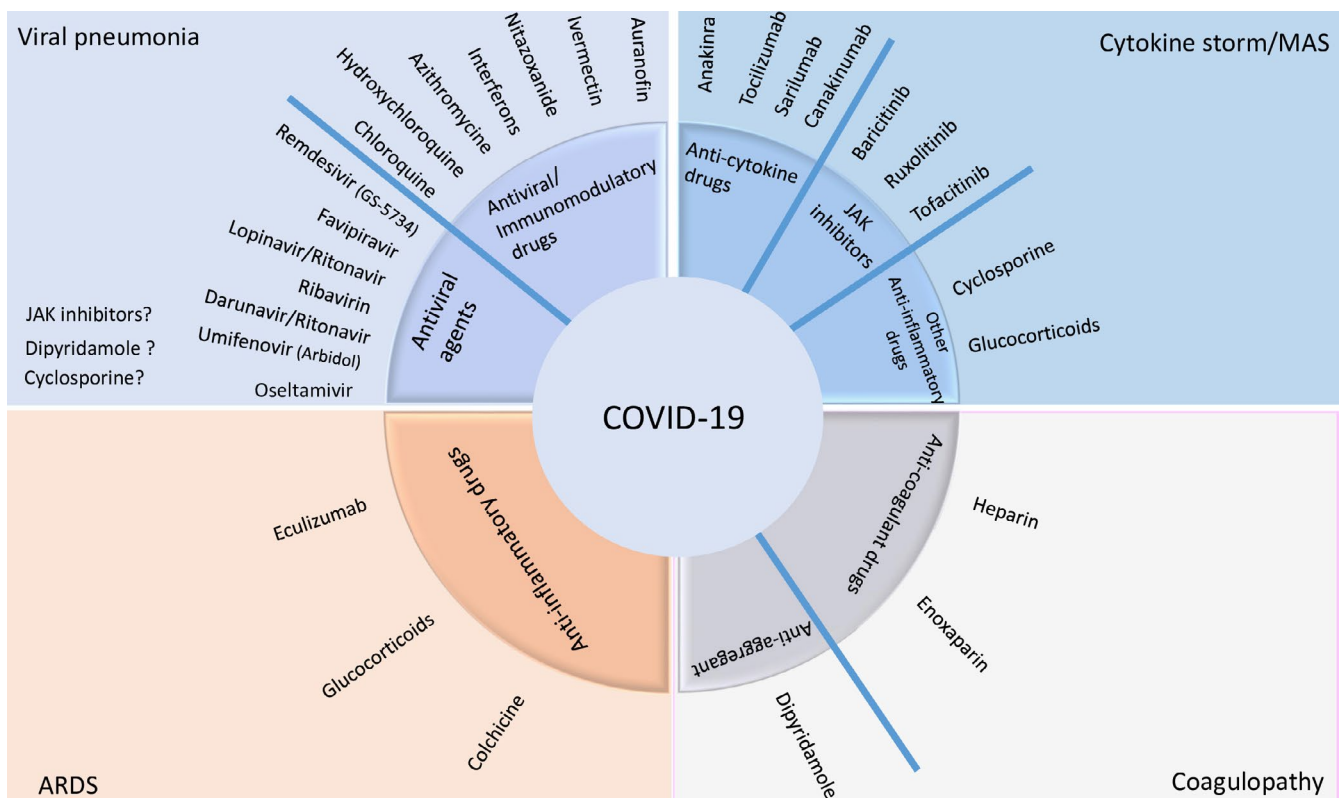


FIGURE 1 Currently investigated drugs in COVID-19 grouped according to their clinical use

2 | SKIN MANIFESTATIONS INDUCED BY COVID-19

There have been increasing reports of dermatologic manifestations associated with COVID-19 (Table 1). It is knowledge, although in progress, rapidly evolving as evidenced by most publications being ahead of print and available only in an electronic version or reported in networks.

According to pathogenetic mechanisms, skin manifestations reported so far can be divided into (1) skin manifestations similar to those in other viral infections and (2) skin manifestations related to thrombovascular events and vascular pathologies.

2.1 | Skin manifestations similar to those in other viral infections

During the COVID-19 outbreak in China, it was not a focus to document skin manifestations. Consequently, skin rash has only been

reported in 2 out of 1.099 infected patients (0.2%).²¹ In contrast, a study by dermatologists from Italy reported skin manifestations in 18/88 patients (20.4%) with COVID-19.⁴ Cutaneous manifestations seen were either erythematous rash ($n = 14$), widespread urticaria ($n = 3$), or chickenpox-like vesicular rash ($n = 1$). In Spain, among 375 patients with suspected or confirmed COVID-19, maculopapular eruptions (MPEs), sometimes similar to pityriasis rosea, were observed in 47% of the cases, urticarial lesions in 19%, and vesicular eruptions of the trunk in 9%.²² Another case of urticaria was presented in France (Figure 2A)²³ and patients with morbilliform exanthem in the USA (Figure 2B).²⁴ Varicella-like lesions predominantly on the trunk were described in 22 patients with proven COVID-19 infection in Italy.²⁵ Predominance of vesicles was reported in 54.5% and generally mild itching in nine (40.9%) patients. The vesiculopapular exanthem appears to develop early in the course of the disease (Figure 2C).^{22,26} An outbreak of severe Kawasaki-like disease has been reported at epicenters of COVID-19 infection also associated with a polymorphic rash in 30%-50% of affected children.^{27,28} In

TABLE 1 Skin manifestations reported associated with COVID-19

Manifestation	Clinical description	Relative frequency*	Similarity to skin rashes of other infections	References
1. Skin manifestations similar to those in other viral infections				
Acute urticaria	Sudden appearance of wheals with a fleeting nature. Continual appearance and disappearance of new lesions is characteristic.	19%	Unspecific for COVID-19; infections are common elicitors for acute urticaria	(4,22,23,31)
Maculopapular exanthem ("erythematous rash")	Acute erupting, widespread distribution of multiple small, round to oval erythematous macules and/or papules with different degrees of confluence. Mostly trunk, low pruritus.	47%	Unspecific for COVID-19; infections are common elicitors for maculopapular exanthem	(4,21,22,24,31,32)
Varicella-like exanthem ("chickenpox-like rash")	Monomorphic papulovesicular skin eruption. Erythematous papules and vesicles bilaterally and symmetrically mostly on the trunk.	9%	May be more specific for COVID-19, vesicles are quite uncommon for virus exanths and more specific for varicella	(4,22,25,26)
Symmetrical intertriginous exanthem	Flexural erythematous maculopapular exanthem on axillary lesions and trunk +/-antecubital fossa.	Individual case reports	Untypical for infectious exanths	(30)
2. Skin manifestations associated with vascular pathologies				
Purpuric exanthem ("purpuric rash")	Skin rash with petechiae.	Individual case reports	Untypical for infectious exanths, except, for example Parvovirus B19	(22,33)
Erythema ab igne ("livedo reticularis")	Transient macular erythema in a broad reticular pattern on thigh unilaterally.	6% together with cutaneous acro-ischemia	Untypical for infectious exanths	(34)
Chilblain-like lesions	Acute-onset, violaceous, infiltrated, and painful plaques on the toes and lateral feet. Vesicles and erosions may be present.	19%	Untypical for infectious exanths	(22,35,36,37)
Cutaneous acro-ischemia	Finger and toe cyanosis, purpura, hematoma, skin bulla, and dry gangrene.	6% together with Erythema ab igne ("livedo reticularis")	Typical for severely ill patients with sepsis	(38,39)

*Relative frequency in percent of this skin manifestations associated with COVID-19 infections according to Ref. (26). In cases, where no numbers are given, only individual case reports do exist.

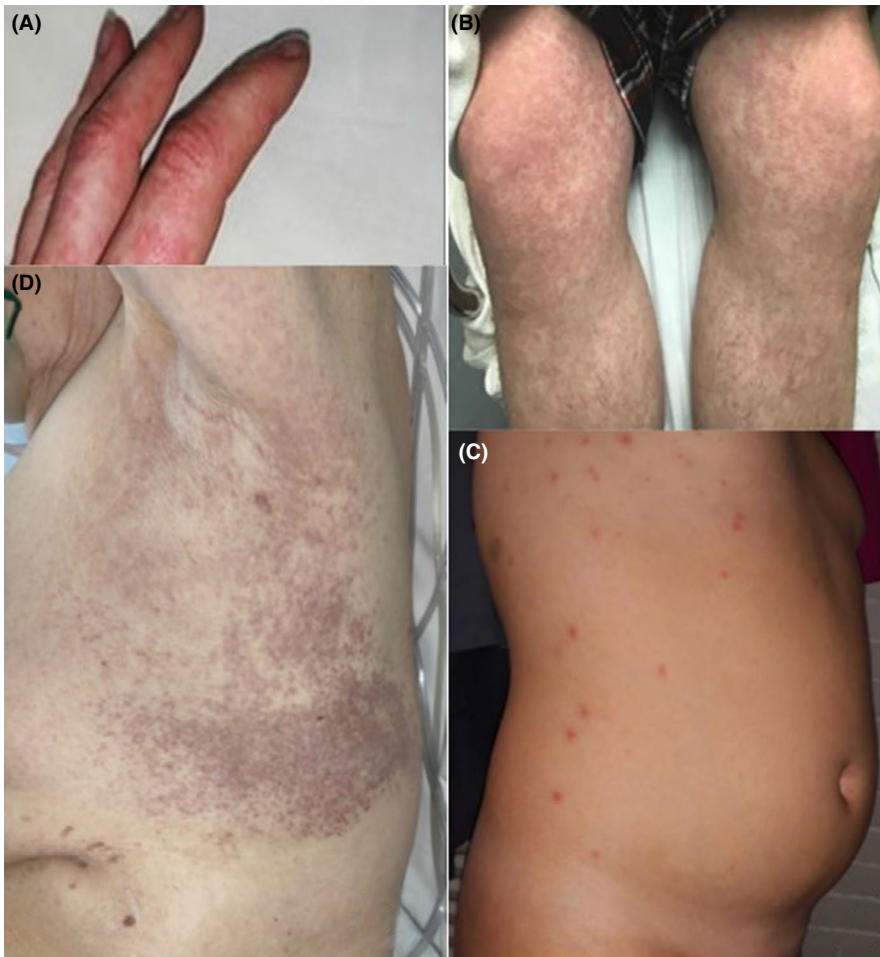


FIGURE 2 Skin manifestations similar to those in other viral infections. A, Urticaria,²³ B, Morbilliform maculopapular exanthem,²⁴ C, Vesiculopapular (chickenpox-like) exanthem,²⁶ D, Intertriginous purpuric rash³⁰

one case, picture of a urticaria-like rash in a 6-month-old child with Kawasaki-like disease associated with COVID-19 infection is shown (Figure S1).²⁹ Two patients with bilateral flexural exanthems resembling systemic drug-related intertriginous exanthems (SDRIFE), one with axillary purpuric lesions associated with thrombocytopenia, have been published (Figure 2D).³⁰ A prospective study from France reported a prevalence of 5/103 (4.9%) and confirmed association of pruritic erythematous rash (n = 2) and urticaria (n = 2) with COVID-19 infections³¹; they additionally observed one oral herpes simplex virus type 1 reactivation. The histopathological picture of exanthematic skin lesions generally resembles that of viral exanthems. However, in individual patients, early microthrombi and an interface dermatitis with necrotic keratinocytes surrounded by lymphocytes have been reported.³²

2.2 | Skin manifestations associated with thrombovascular events and vascular pathologies

COVID-19 exanthems have also been reported with petechiae and low platelet count resembling dengue.³³ In two patients, unilateral lesions on the thigh resembling *livedo reticularis* or erythema *ab igne* have been described with microthromboses discussed as possible etiology (Figure 3A).³⁴

Chilblain-like skin lesions have been frequently reported to be associated with COVID-19^{22,35-37} (Figure 3B).³⁵ They appear in up to 19% of patients, typically in mildly affected ones, and late in the evolution of the disease.^{22,37} Vesicles, pustules, and erosions on these violaceous plaques may occur.³⁷ In Spain, they were observed in 19% of 375 cases.²²

Seven patients had cutaneous acro-ischemia including finger and toe cyanosis, skin bulla, and dry gangrene associated with COVID-19 infection-induced hypercoagulation including definitive DIC in four patients. Five of these patients finally died (Figure 3C).³⁸ A catastrophic microvascular injury syndrome mediated by activation of complement pathways and an associated procoagulant state were described in severe COVID-19 with purpuric skin rash in 3/5 patients.³⁹

In conclusion, the prevalence of cutaneous manifestations in COVID-19-infected patients has been reported between 0.2%, 4.9%, and 20.9%.^{4,21,30} Most skin manifestations resemble cutaneous involvement commonly occurring during viral infections, that is, erythematous rash and acute urticaria. Drug exanthems have to be considered as differential diagnosis.¹⁵ Vesicular varicella-like exanthems may be more specific for COVID-19. Flexural distribution, and petechiae as well as erythema *ab igne*-like lesions have been described. Violaceous, infiltrated painful plaques resembling chilblains have been frequently reported and discussed as typical



FIGURE 3 Skin manifestations associated with thrombovascular events and vascular pathologies. A, Transient unilateral livedo reticularis (erythema ab igne),³⁴ B, COVID-19-induced chilblains,³⁵ C, Acro-ischemia with cyanosis, skin bulla, and dry gangrene in critically ill patient³⁸

manifestations. Necrotic lesions occurred in older and in severely ill patients with increased mortality.²² Cutaneous acro-ischemic microthromboses and small blood vessel occlusion have to be further explored for their causality and specificity for COVID-19 manifestations.

3 | ANTIVIRAL AGENTS USED FOR VIRAL PNEUMONIA

3.1 | Clinical use in COVID-19

Most antiviral agents used for COVID-19 act either by inhibiting RNA-dependent RNA polymerase [remdesivir (GS-5734)] or proteases [lopinavir/ritonavir (LPV/r), favipiravir (FPV), ribavirin, and darunavir].⁴⁰⁻⁴³ Additionally, umifenovir plays a role in viral entry by inhibiting the hemagglutinin-mediated membrane fusion, and oseltamivir is a neuraminidase inhibitor which blocks the release of viral particles from the host cells in influenza infection.⁴⁴ Remdesivir and FPV are considered to be the most effective agents and are mostly used in combination with other COVID-19 medications like hydroxychloroquine.⁴⁰⁻⁴³ Oseltamivir is recommended for concomitant influenza infection.⁴⁵ Darunavir or LPV/r can be concomitantly administered with chloroquine or hydroxychloroquine.⁴³

3.2 | Hypersensitivity reactions

Drug hypersensitivity reactions to ribavirin, darunavir, LPV/r, remdesivir, and oseltamivir are rarely reported, whereas no DHRs to favipiravir and umifenovir are known at present⁴⁶⁻⁵¹ (Table 2).

3.2.1 | Ribavirin

Ribavirin is used in combination with pegylated-interferon $\alpha 2a$ (peg-IFN- $\alpha 2a$) for treating chronic hepatitis C, and both have been

associated with several cutaneous DHRs.⁵² Ribavirin alone causes dermatitis, alopecia, and photoallergic eczematous reactions,^{53,54} and the risk of DHR increases with combination therapy: rash [response rate (RR), 1.74; 95% confidence interval (CI), 1.17-2.6], dermatitis (RR, 1.67; 95% CI, 1.21-2.30), and pruritus (RR, 1.62; 95% CI, 1.29-2.02).⁵⁵ A meta-analysis revealed that, on combination therapy, mild to moderate cutaneous reactions appear in 13.3% of patients, localized cutaneous reactions in 2.6%, generalized reactions-pruritus, skin xerosis, and eczematous changes in 10.3%, alopecia in 4.1%, and exacerbation of lichen planus in less than 1%⁴⁶ (Table 2).

The etiological diagnosis is difficult in case of combination therapy. A drug provocation test (DPT) confirmed the diagnosis of ribavirin hypersensitivity in a patient having MPE due to combined use of peg-IFN- $\alpha 2a$ and ribavirin.⁵⁶ In another case, an erythema multiforme-type drug eruption occurred with peg-IFN- $\alpha 2a$, ribavirin, and/or fluvastatin sodium therapy and a positive lymphocyte transformation test (LTT) confirmed the diagnosis of ribavirin hypersensitivity.⁵⁷ Successful desensitization protocols were reported^{58,59} (Table 2).

3.2.2 | Lopinavir/ritonavir (LPV/r)

Lopinavir/ritonavir, either alone or in combination, has been rarely reported to be associated with DHRs. In human immunodeficiency virus (HIV)-infected patients who received LPV/r combination, MPE rate was reported as 2%-4%.⁶⁰ Acute generalized exanthematous pustulosis (AGEP) was described in two cases receiving LPV/r⁶¹ (Table 2).

In a multicentre randomized study that evaluated the long-term efficacy and safety of the combination of efavirenz or LPV/r plus abacavir/lamivudine, 2/63 patients in the LPV/r group discontinued the study because of a DHR.⁶²

In a recent cohort of 199 severe COVID-19-infected patients who received LPV/r combination, only two (1%) experienced self-limited skin eruptions.⁴⁷ A recent study evaluating 217 patients

TABLE 2 Hypersensitivity reactions due to drugs with antiviral properties investigated for the treatment of COVID-19 in clinical trials or in vitro studies

Drug groups	Drugs	Purpose of use in COVID-19	Hypersensitivity reactions	In vivo tests in IHRs	In vivo tests NIHRs	In vitro tests for IHRs	In vitro tests for NIHRs	Desensitization	
Antiviral drugs	Favipiravir	Viral	None						
	Lopinavir/Ritonavir	pneumonia	AGEP ⁶¹ , MPE ⁶⁰						
	Darunavir/Ritonavir		MPE ⁶⁶ Vesiculobullous lesions ^{48,64}					DNIHR ^{66,67}	
	Umifenovir (Arbidol)		None						
	Ribavirin		Pruritus ^{46,55} Eczema ^{46,53-55} Urticaria ⁵⁸ MPE ⁵⁶ MPE ⁵¹		DPT ⁵⁶	LTT ⁵⁷		DNIHR ^{58,59}	
	Remdesivir (GS-5734)		Anaphylaxis ⁶⁸ SJS/TEN ⁴⁶ , TEN ⁴⁷		SPT ⁶⁸		LTT ⁵⁰		
Immunomodulatory drugs	Azithromycine		MPE ^{73,74} , ACD ^{75,76} , FDE ⁷⁷ AGEP ⁷⁸ , DRESS ⁷⁹ SJS ^{80,81}	SPT ^{84,85} IDT ^{84,85} DPT ⁸⁷	PT ⁷⁵ DPT ⁸⁷			DIHR ⁸⁸	
			Anaphylaxis ⁷² Urticaria ⁷¹ Leukocytoclastic vasculitis ⁸² Hypersensitivity myocarditis ⁸³						
	Hydroxychloroquine / Chloroquine		MPE ⁹⁰ , AGEP ^{92,103} DRESS ⁹³⁻⁹⁵ Erythema multiforme ⁹⁶ Bullous erythema ⁹⁷ SJS/TEN ^{99,100}	SPT ¹⁰⁹	PT ^{96,98,103} DPT ⁹⁰				DNIHR ¹⁰⁴⁻¹⁰⁸ DIHR ^{109,110}
			Photoallergic dermatitis ¹⁰¹ ACD ¹⁰² Anaphylaxis ^{109,110}						
	Auranofin		None						
		Interferons		Local reaction ¹¹⁶ Urticaria ^{119,122,123} Eczema ¹¹⁹ , FDE ¹²⁰ , MPE ¹¹⁹ Anaphylaxis ^{124,125}	SPT ¹²²⁻¹²⁴ DPT ¹²⁵	IDT ^{119,126} PT ¹¹⁹			DIHR ¹²³ DNIHR ^{119,127}
	Nitazoxanide		None						
		Ivermectin		FDE ¹²⁹ , DRESS ¹³⁰ , SJS ¹³¹ , TEN ¹³²					

Abbreviations: ACD, Acute contact dermatitis; AGEP, Acute generalized exanthematous pustulosis; BAT, Basophil activation test; DIHR, Desensitization for immediate hypersensitivity reactions; DNIHR, Desensitization for nonimmediate hypersensitivity reactions; DPT, Drug provocation test; DRESS, Drug-related eosinophilia systemic symptoms; FDE, Fixed drug eruption; IDT, Intradermal test; IHR, Immediate hypersensitivity reaction; LTT, Lymphocyte transformation test; MPE, Maculopapular eruption; NIHR, Nonimmediate hypersensitivity reaction; PT, Patch test; SJS, Stevens Johnson syndrome; SPT, Skin prick test; ST, Skin test; TEN, Toxic epidermal necrolysis.

from China revealed that most of the adverse drug reactions (ADRs) were associated with LPV/r and umifenovir with 63.8% and 18.1%, respectively, and history of a drug allergy was higher in these patients (8.5%) comparing with the ones without ADRs (2.2% vs, $P < .044$).⁶³

3.2.3 | Darunavir

Darunavir can induce a variety of delayed skin eruptions from mild MPE in most cases, to severe bullous cutaneous reactions in HIV-infected patients.^{48,64} A phase III randomized clinical trial performed in 604 patients treated with darunavir/r or LPV/r showed that the percentage of patients experiencing rash was higher in those receiving darunavir/r compared with others (16% vs 7%). Two patients receiving darunavir/r required treatment cessation due to a severe rash.⁴⁸ Darunavir contains a sulfonamide moiety and should be used with caution in patients with a known sulfonamide allergy.⁶⁵ Desensitization was reported to be successful in patients with nonimmediate hypersensitivity reactions (NIHRs) to darunavir^{66,67} (Table 2).

3.2.4 | Oseltamivir

Oseltamivir, used in influenza, causes rare hypersensitivity reactions although close monitoring of patients is important as two cases with Stevens-Johnson syndrome (SJS)/Toxic epidermal necrolysis (TEN) have been reported,^{49,50} with only one being confirmed by LTT.⁵⁰ Another case report revealed anaphylaxis due to oseltamivir confirmed by a skin prick test (SPT)⁶⁸ (Table 2).

3.2.5 | Remdesivir

A recent multicentre study showed that only one (1.6%) out of 61 patients with COVID-19, experienced MPE during remdesivir treatment and therefore discontinued it prematurely⁵¹ (Table 2).

4 | ANTIVIRAL AND/OR IMMUNOMODULATORY DRUGS USED FOR VIRAL PNEUMONIA

4.1 | Azithromycin

4.1.1 | Clinical use in COVID-19

Azithromycin interferes with virus internalization process in influenza infection⁶⁹ and has shown clinical effects in COVID-19-infected patients, although its mechanism against SARS-CoV-2 remains unclear.⁷⁰

4.1.2 | Hypersensitivity reactions

Regarding immediate hypersensitivity reactions (IHRs), urticaria is the most frequent manifestation⁷¹; furthermore, anaphylaxis can occur.⁷² Concerning NIHRs, MPE is described to occur independently⁷³ or only in the presence of a concurrent infection.⁷⁴ Azithromycin has been implicated in contact dermatitis in occupational⁷⁵ and nonoccupational settings.⁷⁶ Cases of fixed drug eruption (FDE),⁷⁷ AGEP,⁷⁸ and Drug Reaction with Eosinophilia and Systemic Symptoms (DRESS),⁷⁹ SJS,^{80,81} leukocytoclastic vasculitis,⁸² and hypersensitivity myocarditis⁸³ were reported (Table 2).

Diagnosis is complex as skin testing is not validated, presenting discrepancies in nonirritating dilutions for SPT and intradermal test (IDT).^{84,85} For NIHRs, positive responses to patch tests (PTs) were described.⁷⁵ In addition, no validated in vitro tests are available.⁸⁶ Oral DPT remains as the gold standard for diagnosis.⁸⁷ A successful desensitization protocol was reported in a case of mast cell activation syndrome⁸⁸ (Table 2).

4.2 | Hydroxychloroquine/Chloroquine

4.2.1 | Clinical use in COVID-19

Hydroxychloroquine/chloroquine have in vitro antiviral effects against SARS-Cov-2 by preventing virus/cell fusion, and immunomodulatory effects by inhibiting production of inflammatory cytokines.⁸⁹

4.2.2 | Hypersensitivity reactions

Dermatologic ADRs are difficult to be distinguished as a side effect of or an allergic reaction to these drugs or a flare of the underlying dermatological disease.^{90,91} The most common manifestation is mild pruritic MPEs within initial 4 weeks of treatment.⁹⁰ High association with AGEP [OR: 39 (8-191)] was described.⁹² Cases of DRESS,^{93,94} pustular DRESS,⁹⁵ erythema multiforme,⁹⁶ bullous erythema,⁹⁷ SJS/TEN,⁹⁸⁻¹⁰⁰ photoallergic dermatitis,¹⁰¹ and occupational contact dermatitis¹⁰² have been reported (Table 2).

PTs are reported to be useful for the diagnosis of NIHRs,^{96,98,103} confirming a T cell-mediated mechanism. However, in a series of 14 patients with ADRs due to chloroquine/hydroxychloroquine, skin tests (STs) were negative in all cases.⁹⁰ DPT is useful in nonsevere cutaneous ADRs in order to differentiate allergic reactions from dermatological adverse effects since only 30% of the patients reporting cutaneous ADRs reveal a positive DPT.⁹⁰ Successful desensitization protocols of hydroxychloroquine in MPE were reported.¹⁰⁴⁻¹⁰⁷ Recently, a 5-hour desensitization protocol for nonimmediate urticaria was successfully administered¹⁰⁸ (Table 2).

Two cases of IHR were reported^{109,110} and one was confirmed by SPTs¹⁰⁹; however, there are no available data for in vitro diagnosis. A hydroxychloroquine desensitization procedure that enables the turning of positive SPTs into negative was published.¹⁰⁹ In a case

of anaphylaxis, a 7 day-desensitization procedure was successfully performed with premedication¹¹⁰ (Table 2).

4.3 | Auranofin

4.3.1 | Clinical use in COVID-19

Auranofin is an anti-inflammatory compound that can possibly inhibit the replication of SARS-CoV-2 in cell culture and reduce the expression of cytokines caused by SARS-CoV-2 and the associated lung damage.¹¹¹

4.3.2 | Hypersensitivity reactions

There are no reported hypersensitivity reactions due to auranofin.

4.4 | Interferons

4.4.1 | Clinical use in COVID-19

Type I IFNs (IFN- α and IFN- β) can inhibit the replication of both SARS and Middle East respiratory syndrome coronavirus (MERS-CoV) and are recommended in combined therapies with other antiviral agents.^{112,113}

4.4.2 | Hypersensitivity reactions

Cutaneous eruptions induced by IFNs are common, with an incidence of 13%-23%.^{114,115} Localized reactions at injection sites are most frequent at 48 weeks.¹¹⁶ Diffuse skin symptoms including urticaria, generalized eczema, papules are common and mostly treated with symptomatic treatment.^{114,117,118} Among 26 patients with non-immediate reactions to IFNs, 12 cases reported generalized eczema, 10 MPE, 3 generalized urticaria, and 1 lichenoid eruption.¹¹⁹ Cases of FDEs,¹²⁰ and subacute cutaneous lupus¹²¹ were described (Table 2).

There are few case reports of immediate urticaria^{122,123} and anaphylaxis.^{124,125} For IHRs to IFN- β , positive STs were reported.^{122,124} For NIHRs, PTs have a low value and are not recommended, whereas delayed reading IDTs are useful.^{119,126} A positive DPT was reported in a patient experiencing anaphylaxis due to peg-INF- α 2a with negative STs.¹²⁵ Successful desensitization protocols both for IHRs¹²³ and NIHRs^{119,127} due to different IFNs were reported (Table 2).

4.5 | Ivermectin

4.5.1 | Clinical use in COVID-19

Ivermectin is an antiparasitic drug also shown to have an in vitro activity against SARS-CoV-2 by inhibition of viral replication.¹²⁸

4.5.2 | Hypersensitivity reactions

Rare case reports of multiple FDEs,¹²⁹ confirmed DRESS by skin biopsy and blood eosinophilia,¹³⁰ confirmed SJS¹³¹ and TEN¹³² by skin biopsy were published (Table 2). No data about STs, in vitro tests, or DPT are available. In addition, no cases of desensitization were reported.

4.6 | Nitazoxanide

4.6.1 | Clinical use in COVID-19

Nitazoxanide is an antiparasitic agent which also has antiviral activities. Combined with hydroxychloroquine or azithromycin, a synergistic effect has been suggested as hydroxychloroquine and azithromycin inhibit viral entry and fusion, while nitazoxanide up-regulates innate immune response to prevent ongoing viral replication in COVID-19.¹³³

4.6.2 | Hypersensitivity reactions

No DHRs to nitazoxanide are reported.

5 | ANTI-CYTOKINE/ANTI-INFLAMMATORY DRUGS USED FOR MAS/CYTOKINE STORM/ARDS

5.1 | Tocilizumab

5.1.1 | Clinical use in COVID-19

Tocilizumab, an anti-IL-6 receptor humanized monoclonal antibody, is under investigation for treatment of COVID-19 and has shown promising results in cytokine storm.⁶

5.1.2 | Hypersensitivity reactions

The rate of all ADRs to tocilizumab is reported to be around 8%, among them 0.1%-0.7% are DHRs.¹³⁴ DHRs to tocilizumab are both NIHRs^{135,136} and IHRs¹³⁷⁻¹⁴⁰ (Table 3). In an adult cohort, the incidence of IHRs was reported as 5.5%¹³⁹ whereas in a pediatric cohort it was 13.6%.¹³⁷

Regarding NIHRs, cases of nonimmediate urticaria,¹⁴¹ DRESS,^{142,143} SJS,¹⁴⁴ and AGEP¹⁴⁵ were reported. Younger age, shorter stature, lighter weight, and increased disease activity in the early period of tocilizumab administration have been identified as risk factors for DHRs.¹⁴⁶

Although not standardized, DPTs, SPTs, and IDTs were used for diagnosis of IHRs in case reports.^{137,139} Only one study

TABLE 3 Hypersensitivity reactions due to other drugs investigated for the treatment of COVID-19–related complications in clinical trials or in vitro studies

Drug groups	Drugs	Purpose of use in COVID-19	Hypersensitivity reactions	In vivo tests in IHRs	In vivo tests in NIHRs	In vitro tests for IHRs	Desensitization	
Anti-cytokine or anti-inflammatory drugs	Tocilizumab	Cytokine storm/MAS	Papular skin lesions ¹³⁶	SPT ^{137,139}	IDT ¹⁴⁰		DIHR ¹³⁴	
			Nonimmediate urticaria ¹⁴¹	IDT ^{137,139}			DNIHR ¹⁴¹	
			Anaphylaxis ^{137,139} DRESS ^{142,143} AGEP ¹⁴⁵ , SJS ¹⁴⁴	DPT ^{137,139}				
	Sarilumab		Pruritic rash ¹⁵⁵					
			ISR ^{147,148} U/Angioedema ¹⁴⁹ Anaphylaxis ^{150,151} Erythematous plaques ¹⁵² U ¹³⁷	SPT ^{150,151} IDT ^{150,151}				DIHR ^{149,151} DNIHR ¹⁵²
	Canakinumab		Palmoplantar pustulosis ¹⁶⁰					
			JAK inhibitors Baricitinib					
			JAK inhibitors Ruxolitinib					
	JAK inhibitors		morbiliform rash, exfoliative dermatitis ¹⁵⁹ U ¹⁶¹ ,					
			Tofacitinib	palmoplantar pustulosis ¹⁶²				
Cyclosporine		Anaphylaxis ¹⁶⁴⁻¹⁶⁷	SPT ^{164,166} IDT ^{164,166}			BAT ¹⁶⁶		
Anti-inflammatory drugs	Glucocorticoids	Cytokine storm/MAS ARDS	IHR ¹⁷⁸⁻¹⁸³	SPT ¹⁷⁸⁻¹⁸⁵	IDT ^{179,189}	slgE ^{181,182}	DIHR ^{186,187}	
			NIHR ^{178,179} ACD ¹⁷⁹	DPT ¹⁸⁰⁻¹⁸⁵	PT ^{179,189} DPT ^{179,189}	BAT ^{181,182}		
			Anaphylaxis ¹⁵¹ FDE ¹⁶⁸ , MPE ¹⁶⁹		DPT ¹⁶⁸ PT ¹⁷⁰			DNIHR ¹⁶⁹
	Colchicine		IHR ¹⁷²⁻¹⁷⁴ Anaphylaxis ¹⁷⁴	SPT ¹⁷⁴ , IDT ¹⁷⁴				DIHR ¹⁷⁴
			ISR ¹⁹² GDE ^{195,196} DRESS ¹⁹⁸ , SJS ¹⁹⁹ HIT ²⁰⁰⁻²⁰² IHR ^{197,204}	SPT ¹⁷ DPT ^{191,192}	PT ¹⁷ DPT ^{191,192} IDT ¹⁷	BAT ^{204,206,207}		DIHR ^{209,210}
	Anti-coagulant or anti-aggregant drugs	Heparin Enoxaparin	Coagulopathy					
				Eczema ²¹⁵ Anaphylaxis ^{216,217}				
Dipyridamole								

Abbreviations: ACD, Acute contact dermatitis; AGEP, Acute generalized exanthematous pustulosis; BAT, Basophil activation n test; DIHR, Desensitization for immediate hypersensitivity reactions; DNIHR, Desensitization for nonimmediate hypersensitivity reactions; DPT, Drug provocation test; DRESS, Drug related eosinophilia systemic symptoms; FDE, Fixed drug eruption; HIT, Heparin induced thrombocytopenia; IDT, Intradermal test; IHR, Immediate hypersensitivity reaction; ISR, Injection site reaction; GDE, Generalized delayed exanthema; LTT, Lymphocyte transformation test; MPE, Maculopapular eruption; NIHR, Nonimmediate hypersensitivity reaction; PT, Patch test; SJS, Stevens Johnson syndrome; SPT, Skin prick test; ST, Skin test; TEN, Toxic epidermal necrolysis; U, Urticaria.

revealed that STs have a low negative predictive value in NIHR.¹⁴⁰ Desensitization to tocilizumab in NIHRs was effectively applied in a weekly scheme with premedication in one case.¹⁴¹ Rapid drug desensitization is successfully and routinely used for IHRs^{19,134} (Table 3).

5.2 | Anakinra

5.2.1 | Clinical use in COVID-19

Anakinra, a recombinant IL-1 receptor antagonist, is under investigation for the treatment of cytokine storm seen during COVID-19.⁵

5.2.2 | Hypersensitivity reactions

Anakinra causes ADRs in 75% of patients. Many of them are related to injection site reactions within the first weeks of application and can present either as an IHR or NIHR.^{147,148} Systemic IHRs such as urticaria, angioedema, anaphylaxis,¹⁴⁹⁻¹⁵¹ and NIHRs¹⁵² as infiltrating erythematous skin plaques were rarely reported as single cases. IHR after a first dose of anakinra was reported in a case possibly due to components that are able to induce a direct mast cell degranulation^{151,153} (Table 3).

For evaluating IHRs to anakinra, SPTs and IDTs were performed with the undiluted drug.^{150,151} For both IHRs^{149,151} and NIHRs,¹⁵² successful desensitization protocols were reported (Table 3).

5.3 | Sarilumab

5.3.1 | Clinical use in COVID-19

Sarilumab, another IL-6 receptor antagonist, is under investigation in a phase II/III clinical trial in patients with severe COVID-19 infection.¹⁵⁴

5.3.2 | Hypersensitivity reactions

It is generally a well-tolerated drug; however, it can cause local reactions on injection site. In an open-label study, in 3% of the patients it caused a pruritic generalized rash which did not affect the treatment¹⁵⁵ (Table 3).

5.4 | Canakinumab

5.4.1 | Clinical use in COVID-19

Canakinumab, a high-affinity human anti-IL-1 β monoclonal antibody, is considered as a candidate in treatment of severe COVID-19.¹⁵⁶

5.4.2 | Hypersensitivity reactions

This anti-IL-1 agent is normally well tolerated and indicated as an alternative in cases with an anaphylactic reaction to anakinra.¹³⁸ However, there is a recently reported case who developed immediate diffuse urticaria after the tenth canakinumab administration and was prevented from further reactions with cetirizine premedication¹³⁷ (Table 3).

5.5 | Janus kinase (JAK) inhibitors (Baricitinib, Ruxolitinib, Tofacitinib)

5.5.1 | Clinical use in COVID-19

Janus kinase inhibitors are under investigation for their potential role in regulating the overactive signaling in the JAK-STAT pathway seen during cytokine storm in critically ill COVID-19-infected patients. Baricitinib with its potential to inhibit clathrin-mediated endocytosis, and its ability to ameliorate associated chronic inflammation in interferonopathies is expected to show promising results in ongoing clinical trials of COVID-19.^{157,158}

5.5.2 | Hypersensitivity reactions

Few cases were reported: one with a morbiliform eruption and exfoliative dermatitis due to ruxolitinib,¹⁵⁹ another one with palmoplantar pustulosis due to baricitinib,¹⁶⁰ and cases of acute urticaria¹⁶¹ and palmoplantar pustulosis¹⁶² due to tofacitinib (Table 3).

5.6 | Cyclosporine

5.6.1 | Clinical use in COVID-19

Cyclosporine A prevents the transcription of genes encoding cytokines like IL-2 and inhibits the replication of diverse coronaviruses at nontoxic, low-micromolar concentrations in vitro.¹⁶³

5.6.2 | Hypersensitivity reactions

Rare cases of pruritus, urticaria, angioedema, and anaphylaxis were reported.¹⁶⁴⁻¹⁶⁶ The possible mechanisms can be both immunologic and nonimmunologic, which seems to depend on the administration route and formulation.¹⁶⁴ In some cases, DHRs have been attributed to the additives such as castor oil¹⁶⁵ or Cremophor EL.¹⁶⁶ SPTs and IDTs or basophil activation test (BAT) can be used for the diagnosis of cyclosporine- and additive-induced IgE-mediated IHRs^{18,164,166} (Table 3).

5.7 | Colchicine

5.7.1 | Clinical use in COVID-19

It is a nonselective inhibitor of NLRP3 inflammasome which is thought to be a major pathophysiologic component of ARDS and/or acute lung injury seen in COVID-19.¹⁶⁷

5.7.2 | Hypersensitivity reactions

Rare cases of anaphylaxis,¹⁵¹ confirmed FDE with DPT¹⁶⁸ and successfully desensitized MPE¹⁶⁹ were reported. For PTs, it is recommended to dilute colchicine to 1% in petrolatum¹⁷⁰ (Table 3).

5.8 | Eculizumab

5.8.1 | Clinical use in COVID-19

Eculizumab, a humanized anti-C5 monoclonal antibody, is under investigation as a candidate drug to play a role in the thrombotic microvascular injury mediated by complement activation causing lung injury either due to severe pneumonia or ARDS in severe COVID-19.^{21,36,171}

5.8.2 | Hypersensitivity reactions

Immediate hypersensitivity reactions or infusion reactions due to eculizumab are very rare.^{172,173} A case of anaphylaxis diagnosed with STs was successfully desensitized with a rapid protocol¹⁷⁴ (Table 3).

5.9 | Glucocorticoids

5.9.1 | Clinical use in COVID-19

In COVID-19-infected patients, the use of glucocorticoids (GCs) is rather controversial.^{175,176} Early start of GCs could be helpful for patients who have an overly exuberant inflammatory response or are at high risk of developing ARDS, whereas the benefit of GCs as rescue treatment remains doubtful.¹⁷⁷

5.9.2 | Hypersensitivity reactions

IHRs to GCs are overall rare and mostly IgE-mediated.¹⁷⁸⁻¹⁸³ In a review of the literature from 2004 to 2014, anaphylaxis was the most common manifestation reported (60.8%, 73/120 reactions) followed by urticaria and/or angioedema (26.7%). Methylprednisolone was implicated in 41% of reactions, followed by prednisolone (20%), triamcinolone (14%), and hydrocortisone (10%).¹⁸¹

In most subjects with IHRs, it is possible to identify the culprit and safe alternative GCs by performing immediate-reading STs.¹⁷⁸⁻¹⁸⁵ In the aforementioned review, 74.1% of 112 STs carried out with GCs suspected of being responsible for reactions were positive.¹⁸¹ In some subjects, positive STs were associated with positive serum-specific IgE assays and BATs^{181,182} (Table 3).

Immediate hypersensitivity reactions to medication components other than the GC itself, such as succinate ester used to enhance the solubility in parenteral preparations, have been described.^{181,185} Hence, when evaluating a reaction to an esterified GC, it is advisable to include in STs the suspected GC and the same GC without the ester component, or with a different ester.

Immediate hypersensitivity reactions to excipients or preservatives in GC preparations, such as lactose, carboxymethylcellulose, polyethylene glycol, and hexylene glycol, have also been reported.^{181,185} Therefore, testing should be performed with a preservative free GC, in addition to preservative testing per se if needed¹⁸⁵ (Table 3). A study proposed a comprehensive diagnostic algorithm to evaluate hypersensitivity reactions to GCs, as well as to their components and preservatives.¹⁸⁵ This algorithm included STs with Carmellose[®] eye drops in subjects who had reacted to carboxymethylcellulose-containing GCs and with cow's milk proteins in those who had reacted to lactose-containing GCs.

In the allergy workup, negative results in STs should be confirmed with DPTs.¹⁸⁰⁻¹⁸⁵ DPTs are also recommended to ensure tolerance of alternative preparations.¹⁸⁴ Cross-reactivity patterns based on structural characteristics have not been clearly established for IHRs as they have been for allergic contact dermatitis.¹⁷⁹ DPTs have shown that patients often tolerate alternative GCs belonging to the same chemical group as the responsible GC.^{182,183} Desensitization to methylprednisolone has been successfully performed^{186,187} (Table 3).

Nonimmediate hypersensitivity reactions following systemic administration of GCs have been more rarely reported than IHRs; most reports concerned isolated cases of eczematous or exanthematous skin eruptions^{178,179} (Table 3). Some are systemic contact dermatitis, occurring in patients with previous contact dermatitis to GCs. They can be revealed by a Baboon syndrome, characterized by a buttock erythema associated to a symmetric, flexural erythema.¹⁸⁸

Most patients do not have a previous topical sensitization. In NIHRs, the main feature is MPE, but other clinical aspects can also occur such as annular erythema, erythroderma, SDRIFE, AGEP, FDE, and a few cases of SJS¹⁸⁸ (Table 3).

Nonimmediate hypersensitivity reactions can be T cell-mediated, and PTs, together with delayed-reading IDTs, are useful tools for evaluating them.¹⁷ PTs have to be read at 2, 4, and also 7 days. Even though delayed reading IDTs are more sensitive than PTs, the sensitivity of the former is limited. Therefore, DPTs are often necessary to diagnose NIHRs. In a study by Padial et al, only 2 of the 38 patients with NIHRs to GCs displayed positive delayed-reading IDTs and PTs to the responsible GCs (ie, dexamethasone and betamethasone), while 21 of the 32 negative patients who agreed to undergo DPTs reacted to them, experiencing almost exclusively delayed-appearing urticarial eruptions or MPEs¹⁸⁹ (Table 3).

6 | ANTI-COAGULANT AND ANTI-AGGREGANT DRUGS USED FOR COAGULOPATHY

6.1 | Heparin and low molecular weight heparins (LMWHs)

6.1.1 | Clinical use in COVID-19

Heparin [unfractionated heparin (UFH)] and LMWHs are administered for treatment or prophylaxis of thrombosis, and therefore, it is used for the coagulopathy observed during COVID-19.¹⁹⁰

6.1.2 | Hypersensitivity reactions

Unfractionated heparin may induce all types of DHRs, mostly type IV and type II.¹⁹¹ Cutaneous NIHRs to subcutaneous heparin occur at the injection site as itchy erythematous or eczematous plaques usually on the 7th-10th day of treatment, although they can appear on the 1-3th day in case of antecedent sensitization.¹⁹² Risk factors for NIHRs to heparin are obesity, female gender, old age, pregnancy, and repeated exposures.^{193,194} If the treatment is continued regardless of a local reaction, the patient may develop generalized eczema or exanthem.^{195,196} Patients with a NIHR to UFH or LMWH at injection site usually tolerate intravenous administration of UFH.¹⁹² Cross-reactivity among LMWHs has been reported in NIHRs.¹⁹⁷ However, fondaparinux is generally well tolerated in patients who react to LMWHs.¹⁹⁴ Heparin may induce DRESS¹⁹⁸ and SJS.¹⁹⁹

Immune-mediated heparin-induced thrombocytopenia (HIT) is induced by IgG antibodies against complex of heparin and platelet-factor 4 tetramers.²⁰⁰ HIT manifests as a more than 50% decrease in the platelet count in 5-10 days after the onset of treatment.²⁰¹ The risk of HIT is increased exclusively with UFH.²⁰² Treatment includes the discontinuation of heparin and the introduction of an alternative anti-coagulant such as argatroban, fondaparinux, danaparoid, or bivalirudin.²⁰³

The IgE-mediated reactions to heparin manifesting as urticaria, angioedema, and anaphylaxis are rare.^{197,203,204} Positive STs with UFH and LMWHs have been reported^{197,203-205} (Table 3). Cross-reactivity in IHRs has been reported between UFH and LMWH and among LMWHs.²⁰⁵

For IHRs with heparins, diagnostic approach primarily consists of SPTs and IDTs.¹⁷ The results of BAT with UFH and LMWH are controversial.^{204,206,207} Heparin itself may cause a release of histamine, leading to a false-positive ST. Further serial dilutions of heparin (1:100, 1:1000, 1:10 000) might be needed.²⁰⁴ IDTs and PTs with the culprit and alternative heparin are performed in NIHRs.¹⁷ PTs, with tape stripping, are less sensitive but may be positive¹⁹¹ (Table 3).

Drug provocation test is considered when the diagnosis is obscure, tissue pathology is unavailable, or an alternative anti-coagulant needs to be determined.²⁰⁸ Subcutaneous DPTs with UFC and LMWHs are performed with increasing doses reaching up to a daily

dose on the first day, then are evaluated on three consecutive days and day 7 in case of NIHRs. Intravenous DPTs with UFC may also be necessary to prove tolerance for emergency situations both for IHRs and NIHRs.^{191,192} A standard protocol for UFH desensitization has not been established yet and published as case reports^{209,210} (Table 3).

6.2 | Dipyridamole

6.2.1 | Clinical use in COVID-19

Dipyridamole is an inhibitor of phosphodiesterase 3 and 5; thereby, it increases intracellular cAMP and/or cGMP in platelets and inhibits platelet aggregation.²¹¹ Besides, it has antiviral features against several viruses.^{212,213} Dipyridamole as an adjunctive therapy was demonstrated to be associated with decreased D-dimer levels in COVID-19.²¹⁴

6.2.2 | Hypersensitivity reactions

Drug hypersensitivity reactions related to dipyridamole are extremely rare. An adult patient with delayed eczematous lesions revealed positive PT results.²¹⁵ Anaphylaxis or anaphylaxis like reactions were described in two cases; however, they lack diagnostic tests^{216,217} (Table 3).

7 | DIAGNOSIS, DIFFERENTIAL DIAGNOSIS, AND MANAGEMENT OF DHRs DUE TO DRUGS INVESTIGATED FOR THE TREATMENT OF COVID-19

Considering the severity of the disease and the emergent need for interventions, it is important to give accurate and quick diagnostic and therapeutic decisions in case of DHRs during COVID-19 treatment. However, it is challenging considering the diverse spectrum of drugs introduced either for direct treatment of the disease or other accompanying conditions during the course of the disease especially in severe cases when the disease is prolonged. Consequently, multiple medications applied at a time make a clear-cut association with one medication more difficult. Furthermore, disease-related eruptions as an important reason of differential diagnosis can make the diagnosis even harder, considering that the majority of the drugs used are more associated with drug-related cutaneous NIHRs.

Given the critical state of the disease, the diagnosis can mostly rely on clinical observations without performing in vivo tests which have possible contamination risks and time-consuming in vitro tests. During a DHR, STs cannot be applied considering the possibility of aggravation and the low diagnostic accuracy expected during ongoing treatment with antihistamines and corticosteroids. When introducing an alternative drug, a DPT based on established methods may be preferred in order to reduce the risk of a possible DHR.¹⁴

BOX 1 Recommendations for diagnosis and management of DHRs in COVID-19

- No equivalent alternatives for the currently off-label repurposed drugs or novel drugs used in COVID-19 do exist.
- We should extrapolate our knowledge on DHRs from other clinical situations to COVID-19 considering the scarce experience for the DHRs during the disease.
- Various drugs being used in different phases of the disease seem to cause rare but potentially severe DHRs, mostly nonimmediate cutaneous hypersensitivity reactions based on data from limited number of case reports.
- The most important differential diagnosis of these DHRs is disease-related exanthems, which can further be classified into the ones similar to those in other viral infections and the others related to thrombovascular events and vascular pathologies seen during COVID-19.
- Experience of diagnostic and management methods for DHRs due to the drugs used in COVID-19 depend mostly on few case reports or series.
- Knowledge of DHRs is urgently needed from pharmacovigilance registries and data from ongoing clinical trials of COVID-19.
- Quick diagnostic and therapeutic decisions in case of DHRs during COVID-19 are mandatory.
- Clinical diagnosis of DHRs during COVID-19 might mostly rely on clinical observations and basic laboratory findings regarding the need of urgent treatment of COVID-19.
- If the risks of a DHR outweigh the benefits obtained from the drug administration, the offending drug should be discontinued.
- When introducing an alternative drug, a DPT may be preferred in order to reduce the risk of a possible DHR.
- If an alternative drug cannot be replaced, the offending drug can be administered via desensitization with published or tailored protocols when there are no contraindications.
- "Treating through" concept, the continued administration of a drug despite a suspected allergic hypersensitivity reaction, can also be considered under strict surveillance measures if the underlying DHR is mild and self-limiting, and an alternative drug does not exist.

If alternative drugs are not available and underlying DHR is not severe, we can recommend that drugs can be applied with published or tailored desensitization protocols.^{19,20} When mild, self-limiting DHR is considered, "treating through" concept, the continued administration of a drug despite a suspected allergic hypersensitivity reaction, can be considered under strict surveillance measures.²¹⁸

Our recommendations for the diagnosis and management of DHRs due to drugs administered during COVID-19 are listed in Box 1.

8 | CONCLUSION

This review brings together all the published information about DHRs due to current and candidate off-label drugs to treat COVID-19. The current knowledge depends mostly on previous clinical experience and few published studies or case reports. Hopefully, published literature reveals that most of these drugs rarely cause DHRs but severe reactions may also occur. One limitation of this review is that it includes extremely low number of reports of ADRs seen so far during COVID-19 treatment. In near future, we need to obtain data about DHRs during the disease from ongoing clinical trials and DHR registries. Additionally as time passes, we will observe if SARS-CoV-2 can aggravate T cell-mediated reactions as some viruses do,²¹⁹ and if the hyperinflammation observed during the course of the disease may influence DHRs.

This review also highlights the presence of two different groups of disease-related exanthems as an important cause of differential diagnosis of DHRs expected during the treatment of the disease. We think that it is extremely important to distinguish these disease-related eruptions from true DHR-related skin manifestations considering that the majority of the drugs used are more associated with drug-related nonimmediate skin reactions.

In near future, further data from ongoing clinical studies and registries established in different countries will enlighten the obscure parts of our understanding on DHRs due to the drugs used in the treatment of COVID-19 and will possibly enable us to establish accurate diagnostic and treatment approaches for these reactions.

CONFLICT OF INTEREST

None of the authors declare conflict of interest.

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SUPPORTING INFORMATION

Additional supporting information may be found online in the Supporting Information section.

How to cite this article: Gelincik A, Brockow K, Çelik GE, et al. Diagnosis and management of the drug hypersensitivity reactions in Coronavirus disease 19: An EAACI Position Paper. *Allergy*. 2020;75:2775-2793. <https://doi.org/10.1111/all.14439>