


Therapeutic Agents Rounding Up the Immunopathology of COVID-19

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Abstract: COVID-19 pandemic has caused more than 3 million deaths globally during the past year. The direct attack from SARS-CoV-2 and hyperactivated immune response contribute to the progress and deterioration of COVID-19. After the virus invades, the activation and release of cytokines/chemokines cause “cytokine storm”, leading to acute respiratory distress syndrome (ARDS) and multiple organs dysfunction syndrome (MODS). Eliminating virus and blocking cytokines are important checkpoints of COVID-19 therapy, and several agents targeting immunopathology, including interferons, thymosin, glucocorticoids and immunoglobulin, have shown therapeutic effects in severe patients with COVID-19. Herein, we reviewed the practice evidences and concluded that several agents rounding up the immunopathology of COVID-19 may be the alternative approaches under the scenario of the lacking of effective antiviral drugs.

Keywords: COVID-19, immunotherapy, cytokines, corticosteroids, inflammation, monoclonal antibodies

Importance

The review provides a summary of the potential drugs related to immunopathology and the practice evidences. Strategies and theoretical bases for the immunological therapy of COVID-19 should be emphasized.

Introduction

The novel coronavirus disease 2019 (COVID-19), caused by acute respiratory syndrome coronavirus 2 (SARS-CoV-2), has been rapidly spreading across the globe despite drastic public health measures in 2020. Siddiqi and Mehra proposed 3 escalating phases of COVID-19 disease progression.¹ During the initial stage, SARS-CoV-2 multiplies and establishes residence in the host, primarily focusing on the respiratory system. Complete blood count may be characterized by lymphopenia and neutrophilia. In the second stage of established pulmonary disease, viral multiplication and localized inflammation in the lung are the norm. A minority of COVID-19 patients will transition into the third and most severe stage of the illness, which manifests as an extrapulmonary systemic hyperinflammation syndrome. In this stage, systemic inflammation markers seem to be elevated. The hyper-inflammation and immune response deregulation contribute to the progress and deterioration of COVID-19 (Figure 1).^{2,3} The lymphopenia is related to delay and decrease of the elimination of virus, and increases the mortality.⁴⁻⁷ The cytokine storm results from a sudden increase of several pro-inflammatory cytokines including IL-6, and IL-1 β , as well as IL-2, IL-8, IL-17, G-CSF, GM-CSF, IP10, MCP1, IFN- γ , TNF- α and

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MIP1 α , etc.⁸ IL-6 is a crucial protagonist for the development of cytokine storm with increased level and lasted for longer duration in severe patients.^{6,9,10} IL-6 further promotes a waterfall-like release of other pro-inflammatory cytokines.^{4,8,11} Besides, monocytes and neutrophils also involve in the cytokine releasing in the severe patients.^{12–14} Eventually, the cytokine storm leads to multiple organs dysfunction syndrome (MODS) and the death (Figure 1).¹⁵ Thus, halting the cytokine storm will be an alternative strategy to improve the outcome for COVID-19. Therefore, we summarized current opinion on the use of several drugs targeting immunopathology in COVID-19 by reviewing the practice evidences available (Table 1), including interferons, thymosin, glucocorticoids (GCs), tocilizumab, and anakinra, etc.¹⁶ In addition, commercial intravenous γ -Immunoglobulin (IVIG), convalescent plasma, and monoclonal antibodies are also investigated for their function in the elimination of virus.^{17,18}

Methods of Selecting Studies for This Review

We conducted a comprehensive literature search in PubMed and EMBASE for English-language articles with available full text that had been published through March 5, 2021. The search terms were combinations of “COVID-19” OR “Coronavirus disease 2019” OR “SARS-CoV-2” AND “Thymosin”, “COVID-19”, OR “SARS-CoV-2” AND “IFN” or “interferon”, “COVID-19” OR “Coronavirus disease 2019” OR “SARS-CoV-2” AND “Glucocorticoids” or “Methylprednisolone” or “Dexamethasone” or “steroid”, “COVID-19” OR “Coronavirus disease 2019” OR “SARS-CoV-2” AND “Tocilizumab” or “TCZ” or “IL-6 receptor Antagonist” or “IL-6 receptor blockade”, “COVID-19” OR “Coronavirus disease 2019” OR “SARS-CoV-2” AND “Anakinra” or “IL-1 receptor blockade” or “IL-1 receptor Antagonist” or “Canakinumab”, “COVID-19” OR

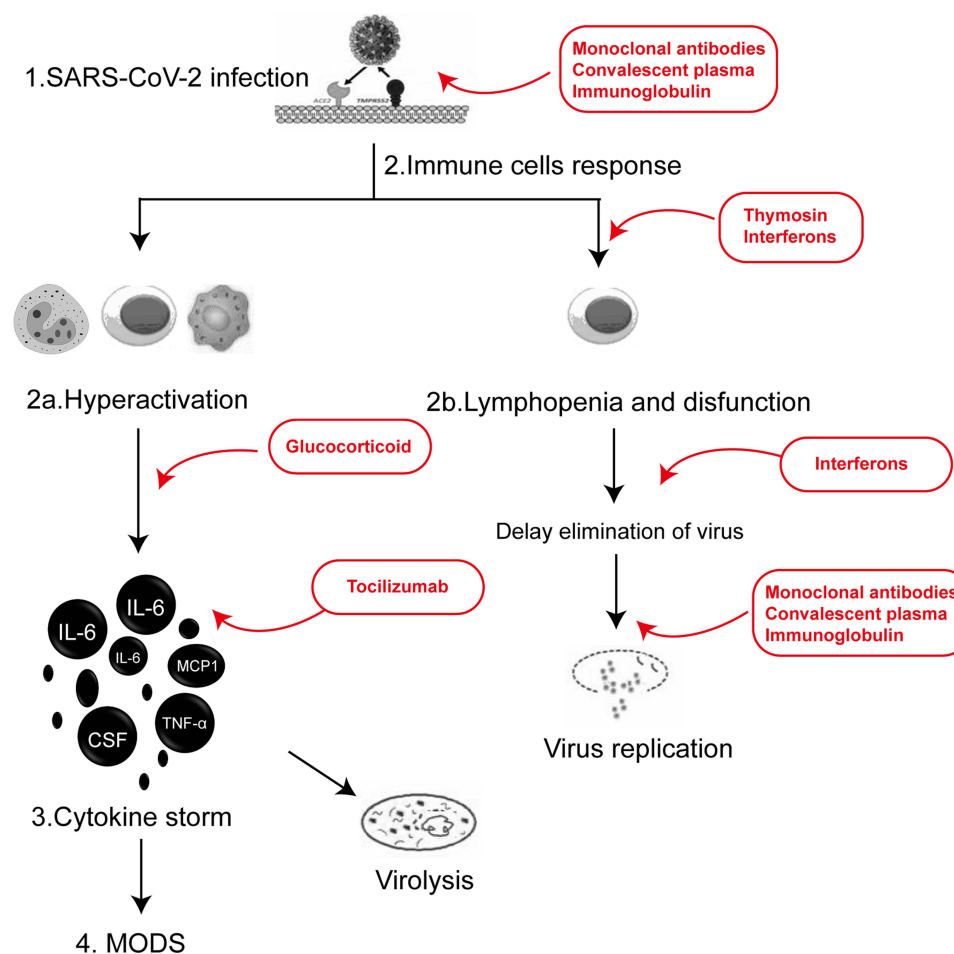


Figure 1 Underlining mechanisms of pathogenesis and potential treatment agents in different steps (numbered) rounding up the immunopathology of COVID-19. 1. Elimination of virus: monoclonal antibodies, inhaling IFN, convalescent plasma and IVIG. 2. Halting hyperreaction and T cell destruction: thymosin (effect of immune regulation and immunosuppression), IVIG and IFN. 3. GC and IL antibodies: inhibiting the cytokines storms.

Abbreviations: IL-6, interleukin-6; IL-1 β , interleukin-1 β ; TNF- α , tumor necrosis factor- α ; MCP1, monocyte chemoattractant protein 1.

Table 1 Clinical Evidence for the Use of Agents Rounding Up the Immunopathology of COVID-19

Agents	Author, Year, Reference	Country	Study Design (Follow-Up), Number	Treatment Regimens	Primary Endpoint	Outcomes
IFN- α -2b	Zhou Q et al, 2020, ³⁰	China	Uncontrolled, exploratory study (Jan. 16 to Feb. 20, 2020), n=77	5 mU b.i.d.	Mean days of detectable virus in the upper respiratory tract	IFN- α -2b: 21.21d ARB+IFN- α -2b: 20.3d ARB: 27.9d
IFN- β -1b+ Ribavirin	Hung IF et al, 2020, ²⁴	China	RCT, prospective (Feb. 10 to Mar. 20, 2020), n=127	8 million IU (three doses on alternate days over a period of 14 days) Ribavirin: 400 mg q12h	The time to providing a nasopharyngeal swab negative for SARS-CoV-2 RT-PCR	IFN- β -1b+Ribavirin: 7d Control: 12d
IFN- β -1a	WHO Solidarity Trial Consortium, 2021, ²⁵	30 countries	RCT (Mar. 22 to Oct. 4, 2020), n=4127 (651: IFN β -1a plus lopinavir)	44 μ g (three doses over a period of 6 days, mainly subcutaneous)	In-hospital mortality	IFN β -1a: 243 of 2050 vs Control: 216 of 2050 (Rate ratio, 1.16; 95% CI, 0.96 to 1.39; P=0.11)
Peg-IFN λ -1	Feld JJ et al, 2021, ²⁶	Canada	RCT (May. 18 to Sept. 4, 2020),	180 μ g (a single subcutaneous injection)	SARS-CoV-2 RNA negative proportion on day 7 after the injection	Peg-IFN λ -1: 24 (80%) vs Control: 19 (63%) (p=0.15)
Thymosin	Wu M et al, 2020, ³⁵	China	Retrospective cohort study (Dec. 2019 to Mar., 2020), n=334	1.6 mg qd, or q12 h for >5 days	28-day and 60-day mortality	28-day mortality Thymosin: 8 of 102 (7.8%) vs Control: 34 of 232 (14.7%) (p=0.016) 60-day mortality Thymosin: 20 of 102 (19.6%) vs Control: 35 of 232 (15.1%) (p=1.000)
Thymosin	Li X et al, 2020, ³⁶	China	Retrospective study (Jan.30 to Apr.2, 2020), n=127	1.6 mg T α 1 (a subcutaneous injection)	Gender differences in CRP, PCT and IL-6 levels	Female: the levels of CRP and IL-6 decreased Males: the decrease of CRP was not significant, and the level of IL-6 was significantly higher than that in females.
Thymosin	Sun Q et al, 2020, ³⁸	China	Retrospective study (Jan.1 to Feb.29, 2020), n=771	Not available	28-day mortality	Thymosin: 8 of 102 (51%) vs Control: 34 of 232 (52.9%) (p>0.05)
Methylprednisolone	Xiang Z et al, 2020, ⁴³	China	Retrospective study, n=9	Body weight \leq 80 kg first 3–4 days: 40 mg/d next 3 days: 20 mg/d Body weight \geq 80 kg first 3–4 days: 80 mg/d next 3 days: 40 mg/d	Clinical parameters: lymphocyte ratio; CRP; IL-6 cytokine levels; chest CT images	Significant improvement: 7 of 9
Dexamethasone	RECOVERY Collaborative Group et al, 2020, ⁴⁴	United Kingdom	RCT (Mar. 19 to Jun. 8, 2020), n=6425	6 mg (oral or intravenous once daily)	28-day mortality	Dexamethasone: 482 of 2104 (22.9%) vs Control: 1110 of 4321 (25.7%) (p<0.001)

(Continued)

Table I (Continued).

Agents	Author, Year, Reference	Country	Study Design (Follow-Up), Number	Treatment Regimens	Primary Endpoint	Outcomes
Methylprednisolone Dexamethasone	Ko JJ et al, 2021, ⁴⁵	USA	Cohort study (Mar. 1 to Jul. 31), n=262	Methylprednisolone 1 mg/kg/day for \geq 3 days Dexamethasone 6 mg/day for \geq 7 days	Mortality within 50 days	Methylprednisolone: 16.4% Dexamethasone: 26.5% vs Control: 41.3% (p<0.001)
Tocilizumab	Luo P et al, 2020, ⁵²	China	Retrospective study (Jan. 27 to Mar. 5, 2020), n=15	80 to 600 mg per time	CRP IL-6 level	126.9 (10.7-257.9) to 11.2 (0.02-113.7) mg/L (p<0.01) In 10 (66.7%) patients decreased
Tocilizumab	Toniati P et al, 2020 ⁵³	Italy	Prospective study (Mar. 9 to Mar. 20), n=100	8 mg/kg (max 800 mg) by two consecutive intravenous infusions 12 h apart.	Lymphocyte count; levels of CRP, fibrinogen, ferritin and IL-6; respiratory condition	Increased Decreased improved or stabilized in 77 (77%) patients
Tocilizumab	REMAP-CAP Investigators et al, 2021 ⁵⁴	NA	RCT (Mar.9 to Jun. 17), n=803	8 mg/kg	Respiratory and cardiovascular organ support-free days	Tocilizumab: 10 (-1 to 16) vs Control: 0 (-1 to 15)
Tocilizumab	Perrone F et al, 2020, ⁵⁵	Italy	Single-arm hypothesis -driven phase 2 study (Mar. 19 to 20, 2020), n=301 Validation cohort (Mar. 20 to 24, 2020), n=920	8 mg/kg to a maximum of 800 mg per dose	Lethality rates at 14 and 30 days (expected lethality rates at 14 and 30 days: 2% and 35%)	Phase 2: 14-day: 55 of 299 (18.4%) 30-day: 67 of 299 (22.4%) Validation 14-day: 101 of 884 (11.4%) 30-day: 158 of 858 (18.4%)
Tocilizumab	Stone JH et al, 2020 ⁵⁷	USA	RCT (Apr. 20 to Jun. 15), n=243	8 mg/kg to a maximum of 800 mg	Intubation or 28-day mortality	Tocilizumab: 17 of 161 (10.6%) vs Placebo: 10 of 81 (12.5%)
Tocilizumab	Veiga VC et al, 2021, ⁵⁸	Brazil	RCT (May. 8 to Jul. 17, 2020), n=129	8 mg/kg (single intravenous infusion)	Death or mechanical ventilation at 15 days	Tocilizumab: 18 of 65 (28%) vs Control: 13 of 64 (20%)
Anakinra	CORIMUNO-19 Collaborative group et al, 2021, ⁶⁰	France	Bayesian RCT (Apr. 8 to Apr. 26, 2020), n=116	200 mg twice a day on days 1-3, 100 mg twice on day 4, 100 mg once on day 5 IL-6 inhibitors (Tocilizumab/Sarilumab): a single dose of 400 mg	A score of >5 on the WHO-CPS at day 4 Need ventilation or died at day 14	A score of >5 on the WHO-CPS at day 4 Anakinra: 21 of 59 (36%) Control: 21 of 55 (38%) Need ventilation or died at day 14 Anakinra: 28 of 59 (47%) Control: 28 of 55 (51%)
IL receptor blockade	Cavalli G et al, 2021, ⁶¹	Italy	Cohort study (Feb. 25 to May. 20, 2020), n=392	Anakinra: 5 mg/kg twice daily Tocilizumab/ Sarilumab: a single dose of 400 mg	Survival	Anakinra vs Control 74 of 100 (86%) vs.61 of 75 (68%) (p=0.047) IL-6 inhibitors vs Control 69 of 97 (82%) vs.61 of 75 (68%) (p=0.79)

(Continued)

Table I (Continued).

Agents	Author, Year, Reference	Country	Study Design (Follow-Up), Number	Treatment Regimens	Primary Endpoint	Outcomes
Anakinra	Cauchois R et al, 2020, ⁶²	France	Retrospective study (20d), n=22	A single dose of 300 mg/d for 5 d; 200 mg/d for 2 d; 100 mg/d for 1 d	NEWS score (severity) at D5 Days with O ₂ < 3L/min ICU free days Invasive ventilation free days Death, n (%)	Anakinra vs Control 4.5 vs.8 (p<0.01) 15.5 vs.8 (p<0.05) 17.5 vs.14 20 vs.17 0 (0%) vs.1 (10%)
Anakinra	Cavalli G et al, 2020, ⁶³	Italy	Retrospective cohort study (21d), n=52	High-dose: 5 mg/kg twice daily	Survival MV-free survival Respiratory function improvement	Anakinra(n=29) vs Control (n=16) 90% vs.56% (p=0.009) 72% vs.50% (p=0.15) 21 (72%) vs.8 (50%)
Anakinra	Huet T et al, 2020, ⁶⁴	France	Cohort study (Mar. 24 to Apr. 6, 2020), n=96	100 mg twice a day for 3d 100 mg daily for 7 days	Admission to the ICU for IMV or death	Anakinra group: 13 of 52 (25%) Historical group: 32 of 44 (73%)
Canakinumab	Landi L et al, 2020, ⁶⁵	Italy	Prospective cohort study (30d), n=88	300 mg sc	Median PaO ₂ /FiO ₂ Median duration of hospitalization Death at 7 days Overall survival at 1 month	From 160 at baseline to 237 6d 12 of 88 (13.6%) 79.5%
Canakinumab	Generali D et al, 2020, ⁶⁶	Italy	Prospective case-control study (01–25 Apr. 2020), n=48	150mg on day 1 and day 7	Survival rate at 60 days	Canakinumab: 90.0% Control: 73.3%
IVIG	Xie Y et al, 2020, ⁶⁸	China	Retrospective study (Jan. to Feb. 2020), n=58	20 g/day	28-day mortality	≤48 h group 7 of 30 (23.3%) >48 h group 16 of 28 (57.1%) (p = 0.009)
IVIG Infiximab	Farrokhpour M et al, 2021, ⁶⁹	Iran	Cohort study (Mar, 2020), n=104	IVIG: 400mg/kg/d for 3 to 5 days Infiximab: a single dose of 5 mg/kg	Death	IVIG: 6 of 23(26.1%) Infiximab: 10 of 27(37%) Combination Therapy: 5 of 11 (45.5%) vs Control: 27 of 43(62.8%) (all P < 0.05)
IVIG	R S R et al, 2021, ⁷⁰	India	Phase-II RCT (Jul. to Sept., 2020), n=100	0.4g/kg/d for 5 days	Number of days of hospitalization	IVIG: 7.72 vs Control: 17.5 (p < 0.01)
IVIG	Huang C et al, 2021, ⁷¹	China	Retrospective Cohort Study (Jan.20 to Jun.10, 2020), n=639	10 g/day for 3/5 days, 20 g/day for 3/5 days,	Severity rate Mortality rate	IVIG: 3 of 45 (6.6%) vs Control: 3 of 90 (3.3%) (p>0.05) IVIG: 1 of 45 (2.2%) vs Control: 0 of 90 (p>0.05)
Convalescent plasma	Duan K et al, 2020, ⁷⁴	China	Retrospective study (Jan. 23 to Feb. 19, 2020), n=10		Improvement of clinical symptoms: fever, cough, shortness of breath, and chest pain; Reduction of pulmonary lesions on chest CT examinations	10 patients different degrees of absorption in 10 patients

Abbreviations: RCT, randomized controlled trial; IL-6, interleukin-6; ICU, intensive care unit; CRP, C-reactive protein; IVIG, intravenous γ -immunoglobulin.

“Coronavirus disease 2019” OR “SARS-CoV-2” AND “Intravenous γ -Immunoglobulin” or “IVIG”, as well as combinations of “COVID-19” OR “Coronavirus disease 2019” OR “SARS-CoV-2” AND “convalescent plasma”. Meta-analyses, case report and review articles were not included but were screened for additional studies that were not found in the initial electronic search. Titles and abstracts were evaluated in order to determine the applicability of each individual study. Finally, 28 papers were retained for this review.

Target Therapeutic Agents Rounding Up the Immunopathology Interferons

Type I interferons (IFNs), including IFN- α and IFN- β , are highly active and multifunctional secreted glycoproteins with antiviral, antitumor, and immunomodulatory functions,¹⁹ which regulate the expression of a series of genes, through binding of IFNs to specific receptors located on the cell membrane. IFNs exert antiviral effect by inhibiting virus replication and regulating immune function.²⁰ McNab has found the deficiency of type-I IFNs might be an indicator of severe COVID-19,²¹ for no IFN- β and low IFN- α production and activity were found in the severe and critical patients, which were associated with a persistent blood viral load and an exacerbated inflammatory response. IFNs help to eliminate the SARS-CoV-2 by improving the phagocytic function of macrophages and the activity of T lymphocytes, or by inducing the synthesis of antiviral protein (AVP) of host cells. Thus, type I IFNs have emerged as potential drugs against SARS-CoV-2.²² Zhou et al found that IFN- α 2b, with or without oral umifenovir hydrochloride, reduced the duration of detectable virus in the upper airway significantly in 53 moderate COVID-19 patients than those who used umifenovir hydrochloride only (21.1 days vs 27.9 days).²³ Over the time period from day 12 (the onset of symptoms) to day 42, patients treated with IFN- α 2b had lower IL-6 levels (33.5 pg/mL) and higher CRP levels (25.7 mg/L) than those without IFN- α 2b treatment. A multicenter study reported that the antiviral therapy combined with IFN- β -1b (8 million IU on alternate days, iH) had a significantly shorter median time from start of treatment to negative nasopharyngeal swab virus test [7 days (IQR 5–11)] than the control group [12 days (8–15); HR 4.37 (95% CI 1.86–10.24), $p = 0.0010$], and days of hospital stay in 86 patients with mild to moderate COVID-19 [9.0 days (7.0–13.0) vs 14.5 days (9.3–16.0); HR 2.72 (1.2–6.13), $p = 0.016$].²⁴ As for the mortality, unfortunately, an interim WHO solidarity trial results did not show that IFN- β -1

definitely reduced initiation of ventilation or hospitalization duration, and the mortality either (Death occurred in 243/2050 with interferon vs 216/2050 in control, $P=0.11$).²⁵

IFN III (IFN- λ) imposes the antivirus effects via binding and inducing signaling through the heterodimeric IFN- λ receptor (IFNLR). A recent Phase 2, placebo-controlled randomized trial reported that Peg-IFN λ -1 has potential to prevent clinical deterioration and shorten duration of viral shedding. Peg-IFN λ -1 accelerated viral decline in outpatients with COVID-19, increasing the proportion of patients with viral clearance by day 7, particularly in those with high baseline viral load.²⁶ However, it should be mentioned that IFN- λ has been shown to increase susceptibility to pneumonia caused by methicillin-resistant staphylococcus aureus in response to influenza virus infection in mice.²⁷ Broggi et al also found that IFN- λ induced barrier damage of airway epithelium in mice.²⁸ The amounts of IFN- λ mRNA in bronchoalveolar lavage fluid and naso-oropharyngeal samples were correlated with the morbidity of SARS-CoV-2-positive patients. Thus, it is cautious to use the IFN- λ .

Being different with IFN I and IFN III, IFN II (IFN- γ) is an independent risk factor associated with mortality in patients with moderate and severe COVID-19, as the higher level of IFN- γ was found in the death than the that of survivals (median of 323 pg/mL vs 208 pg/mL, $p = 0.017$).²⁹

Thymosin

Thymosin, a small molecular peptide with physiological activity is extracted from thymus of animals and human.³⁰ Thymosin alpha 1 is produced as synthetic peptide and the most used form as drug. Thymosin preparations play a major role in maintaining the immune response of patients with primary immunodeficiency, autoimmune diseases and cancer, by stimulating the activity of specific lymphocytes.³¹ Acting as an immune potentiator, thymosin promotes the differentiation and maturation of thymocytes, increases the number and activity of T cells.³² Thymosin produce various cytokines such as IFN- γ , IL-2 and IL-3 through Th1, and increase the expression of IL-2 receptor through the activation of mitogen or antigen.³³ It can also promote the recruitment of precursor NK cells, and the differentiation and function enhancement of bone marrow-derived dendritic cells (DC).³⁴ Activated macrophages or DC secrete IL-12, which strongly induces the differentiation of CD4+ T cells into Th1 cells. Th1 response is accompanied by a significant antiviral response, while Th2 response is accompanied by the continuation of viral

infection and immune escape. In addition, because of its ability to increase the production of macrophage migration inhibitory factor and IFN- γ , it has been widely used for antiviral treatment of hepatitis B and HIV.³²

As for the COVID-19, thymosin might restrict the progression of COVID-19 by stimulating the production and activity of lymphocytes, especially for the patients with lymphopenia. A multicenter retrospective cohort study (334 cases of COVID-19 enrolled) showed that the administration of thymosin- α (1.6 mg qd or q12 h for >5 days) significantly reduced the 28-day mortality (HR, 0.11, $P=0.013$) of the critical patients with lymphocyte $< 0.73 \times 10^9/L$, PaO₂/FiO₂ < 196 , SOFA > 3 , and acute physiology and chronic health evaluation (APACHE) II > 7 .³⁵ Gender differences may be a factor in sustaining COVID-19 immunity responded to thymosin α 1. It was found that the levels of CRP and IL-6 were higher in T α 1-treated male than T α 1-treated female, while the level of PCT was significantly lower in T α 1-treated male group.³⁶ Another study demonstrated that T α 1 mitigates peripheral blood cytokine expression and inhibits lymphocyte activation specifically in a CD8⁺ T-cell subset from patients with COVID-19 in vitro, suggesting the potential to modulate immune response homeostasis and the cytokine storm in vivo.³⁷ However, another multicenter retrospective study (771 cases) did not find the association between the use of thymosin- α 1 and the decreased mortality in critically ill COVID-19 patients.³⁸ The inconsistency may attribute to the difference of study subjects, severity of the COVID-19, proportion of the genders, and the dose and timepoint of administration, etc. Nevertheless, the use of thymosin- α 1 is still controversial.

Glucocorticoids (GCs)

Glucocorticoids (GCs) potently suppress the levels of pro-inflammatory cytokines.³⁹ GCs can inhibit the phospholipase A2, affect a cascading group of reactions of arachidonic acid, stabilize the lysosomal membrane and reduce tissue and cell damage. GCs suppress the binding of NF- κ B to GCs response elements on DNA directly, reducing the expression of cytokines like IL-6, TNF- α , and IL-1 β , etc. It can also activate the inhibitory nuclear factor I κ B α , initiating the transcription of anti-inflammatory factor IL-10.⁴⁰ GCs reduce the permeability of capillaries and improve the sensitivity of blood vessels to catecholamines to constrict blood vessels and reduce inflammation,⁴¹ inhibit the inflammation process of ARDS.⁴² Medium-to-low-dose glucocorticoids (Methylprednisolone 20–40mg/d, less than 8 days) plays

a protective role in the respiratory and digestive systems by activating ACE2 and decreasing the level and IL-6 in 9 severe to critical patients with COVID-19.⁴³ A recent randomized clinical trial showed that the dexamethasone reduced 28-day mortality among the patients with COVID-19, who were treated by invasive mechanical ventilation (IMV) or oxygen at randomization.⁴⁴ In this study, 2104 patients received dexamethasone (6 mg, once daily, oral or intravenous for up to 10 days) reduced deaths by one-third in patients receiving IMV (29.0% vs 40.7%), by one-fifth in patients receiving oxygen without IMV (21.5% vs 25.0%), but did not reduce the mortality of patients without respiratory support at randomization (17.0% vs 13.2%, $p=0.14$), compared with 4321 patients concurrently allocated to usual care. Another study retrospectively compared the effectiveness of methylprednisolone to dexamethasone in patients with COVID-19 requiring intensive care in urban Los Angeles. In this study, all-cause mortalities in the usual care, methylprednisolone, and dexamethasone groups were 41.3%, 16.4% and 26.5% at 50 days ($P < 0.01$) respectively.⁴⁵ It was found that in patients requiring mechanical ventilation, mortality was 42% lower in the methylprednisolone group than in the dexamethasone group (hazard ratio 0.48, 95% CI: 0.235–0.956, $P = 0.0385$).

However, the immunosuppressed effect of GCs should be considered, for the risk of delaying virus clearance and increasing the secondary infection. Thus, the optimal GCs administration juncture timepoint, dosage, and duration might be subjected to further exploration to prevent the progression of ARDS. GCs used in the early stage of the disease did not show significant efficacy, but several studies have confirmed the efficacy of GCs in severe patients.¹⁶ The usefulness of GCs is recommended when the following clinical indicators occur, increased resting respiratory rate (>30 breaths/minute), ongoing deterioration in oxygenation index, multi-lobular progression ($>50\%$) on imaging within 48 h, consistent lymphopenia, etc.^{46,47}

IL-6 Receptor Antagonist

Tocilizumab (TCZ), a humanized IgG1k monoclonal antibody, binds soluble or membrane-type IL-6 receptors (sIL-6R and mIL-6R) specifically, and inhibit the signal transduction mediated by sIL-6R and mIL-6R. In the process of inflammation response, IL-6 regulates innate immunity via interacting with innate immune cells directly or indirectly activating stromal cells on inflammatory sites.⁴⁸ These actions affect the recruitment, functional activation and differentiation of leukocytes, as well as the development of cytokine storm.⁴⁹

TCZ has been widely used in the treatment of autoimmune diseases such as rheumatoid. It was found to reduce the 28-day mortality of patients with sepsis (IL-6 > 1000 pg/mL).⁵⁰

Excessive inflammation response can trigger cytokine storms and cause damage to multiple target organs in COVID-19.¹² IL-6 played a major role in the development of cytokine storm and indicated the poor outcome of severe patients with COVID-19. TCZ was recommended to the patients of COVID-19 with elevated IL-6 levels to suppress the cytokine storm.¹⁰ A case report showed that in a 42-year-old male suffering from respiratory failure due to SARS-CoV-2 infection, after 4 days of TCZ treatment, the CRP decreased from 225 to 33 mg/L and ultimately clinically fully recovered.⁵¹ A retrospective study analyzing 15 cases of COVID-19 by Luo et al found that TCZ decreased the plasma level of CRP and IL-6 in 10 patients.⁵² A single center study of 100 COVID-19 patients with hyperinflammatory syndrome and ARDS demonstrates that ten-day treatment with TCZ increases lymphocyte count, decrease the levels of CRP, fibrinogen, ferritin and IL-6. The study also showed that the respiratory condition was improved or stabilized in 77 (77%) patients.⁵³ An analysis of 90-day survival showed improved survival in the pooled interleukin-6 receptor antagonist groups, yielding a hazard ratio for the comparison with the control group of 1.61 (95% credible interval, 1.25 to 2.08) and a posterior probability of superiority of more than 99.9% for the critically ill Patients.⁵⁴ A multicenter, single-arm, hypothesis-driven trial found that TCZ reduced lethality rate at 14 and 30 days (18.4% vs 20%, $P = 0.52$; and 22.4% vs 35%, $P < 0.001$).⁵⁵ Therefore, the efficacy of TCZ could be expected in severe and critical patients. However, a cautionary case report by Radbel et al demonstrated that two patients with COVID-19 complicated by cytokines release syndrome (CRS) and treated with TCZ, but unfortunately, both patients progressed to severe hemophagocytic histiocytosis (HLH), and one developed to viral myocarditis.⁵⁶ The hazard ratio for intubation or death in the tocilizumab group as compared with the placebo group was 0.83 (95% confidence interval [CI], 0.38 to 1.81; $P = 0.64$), and the hazard ratio for disease worsening was 1.11 (95% CI, 0.59 to 2.10; $P = 0.73$).⁵⁷ In addition, a randomized, open label trial in Brazil found TCZ plus standard care was not superior to standard care alone in improving clinical outcomes at 15 days, and it might increase mortality in patients with severe or critical COVID-19.⁵⁸ Furthermore, a meta-analysis showed that the TCZ did not reduce short-term mortality.⁵⁹ The effect of single anticytokine approach

such as IL-6 receptor antagonist may be limited, possibly because it cannot overcome the complicated inflammatory response in COVID-19.

IL-1 Receptor Antagonist

IL-1 β is also a pro-inflammatory cytokine involved in the inflammatory response to COVID-19, which mediates immune responses and maintains the acute and chronic inflammatory. Early blockade of the IL-1 receptor is therapeutic in acute hyperinflammatory respiratory failure in COVID-19 patients.

Anakinra is an IL-1 receptor antagonist that blocks the activity of IL-1 α and IL-1 β . Recently, a multicenter, open-label, Bayesian randomized clinical trial reported that 21 (36%) of 59 patients in the anakinra group had a WHO-CPS score of more than 5 at day 4 versus 21 (38%) of 55 in the usual care group. At day 14, 28 (47%; 95% CI 33 to 59) patients in the anakinra group and 28 (51%; 95% CI 36 to 62) in the usual care group needed ventilation or died.⁶⁰ These data suggested anakinra did not improve outcomes in patients with mild-to-moderate COVID-19. A cohort study was conducted in Italy to compare the inhibitory effects of IL-1 and IL-6 in patients with COVID-19, respiratory insufficiency, and hyperinflammation. In the multivariable analysis, compared with patients who did not receive IL inhibitors, patients treated with IL-1 inhibition had a significantly reduced mortality risk (hazard ratio [HR] 0.450, 95% CI 0.204–0.990, $p=0.047$).⁶¹ A retrospective study found that all of the patients treated with anakinra improved clinically ($P < 0.01$), with no deaths, significant decreases in oxygen requirements ($P < 0.05$), and more days without invasive mechanical ventilation ($P < 0.06$).⁶² After 21 days treatment with high-dose intravenous anakinra, 21 (72%) of 29 patients with COVID-19, ARDS and hyperinflammation showed reductions in serum C-reactive protein and progressive improvements in respiratory function.⁶³ Thomas et al observed a significant reduction of mortality, along with a significant decrease in the need of mechanical invasive ventilation with no obvious adverse events in severe forms of COVID-19 patients treated with anakinra.⁶⁴ These data suggested that administration of anakinra in patients with an exacerbating form of COVID-19 resulting from cytokine storm has a positive effect.

Canakinumab is a fully human IgG monoclonal antibody that specifically targets IL-1 β . In the US and EU, it is approved for a variety of autoinflammatory syndromes.⁶⁵ An observational, cohort, prospective study showed that patients hospitalized treated with canakinumab, oxygen-

support requirements improved and overall mortality decreased to 13.6%.⁶⁵ A prospective case-control study showed that patients treated with canakinumab experienced a significant increase in PaO₂:FiO₂ ($p < 0.001$) and reduction in lung damage by CT ($p = 0.01$), along with significant decreases in immune/inflammation markers. Thus, canakinumab reduces the need for invasive mechanical ventilation and improve clinical symptoms in patients with moderate COVID-19.⁶⁶

Intravenous γ -Immunoglobulin (IVIG)

The commercial intravenous γ -immunoglobulin (IVIG) is a plasma product extracted from a large amount of healthy human blood. It usually contains main IgG, with little IgA and IgM. IVIG exerts a non-specific antiviral effect and been used as an adjunctive treatment of severe pneumonia related to influenza.⁶⁷ In critically ill patients with COVID-19, the innate immunity is insufficient to limit the virus, and the adaptive immune does not establish usually until 10 days after the onset. Thus, injection of IVIG helps to eliminate the virus partially. Xie et al investigated 58 cases of severe to critical COVID-19 with lymphocyte less than $0.5 \times 10^9/L$, who received IVIG at 20 g/day.⁶⁸ The results showed that IVIG injection within 48 h of admission reduced the proportion of patients requiring mechanical ventilation (6.67% vs 32.14%, $p = 0.016$), length of stay in the hospital (11.50 ± 1.030 vs 16.96 ± 1.620 days, $p = 0.0055$) and ICU stay (11.50 ± 1.030 vs 16.96 ± 1.620 days, $p = 0.0055$), and ultimately decrease the 28-day mortality (23.3% vs 57.1%, $p = 0.009$), comparing with the control. Recently, an observational research investigated the therapeutic effects of infliximab, IVIG, or combination therapy in patients with severe COVID-19 disease admitted to the ICU. According to the findings of this study, the expired rate was 37% in the infliximab group, 26.1% in IVIG, 45.5% in combination therapy, and 62.8% in the control group, respectively (all $P < 0.05$). Hazard ratios of death were 0.31 in IVIG (95% CI: 0.12–0.76, $P = 0.01$), 0.30 in infliximab (95% CI: 0.13–0.67, $P = 0.004$), 0.39 in combination therapy (95% CI: 0.12–1.09, $P = 0.071$).⁶⁹ It seems that infliximab and IVIG, alone or together, in patients with severe COVID-19 disease can be considered an effective treatment. An open-label, multicenter, comparative, randomized study found that duration of hospital stay was significantly shorter in IVIG group than that of standard care alone (7.7 vs. 17.5 days).⁷⁰ Duration for normalization of body temperature, oxygen saturation and mechanical ventilation were significantly shorter in IVIG compared to standard care. However, a retrospective study

for non-severe patients with COVID-19 reported that no statistically significant differences were found between IVIG group and control group in the duration of fever (median, 3 vs 3 days, $p = 0.667$), virus clearance time (median, 11 vs. 10 days, $p = 0.288$), length of hospital stay (median, 14 vs 13 days, $p = 0.469$), and the use of antibiotics (40% vs 38.9%, $p = 0.901$).⁷¹ Thus, IVIG can be considered for use in severe and critically ill patients, although it may not be suitable for non-severe ones. In summary, IVIG acts as a safe and efficacious combination therapy with other antiviral drugs for severe patients with COVID-19.

Convalescent Plasma and Monoclonal Antibody of SARS-CoV-2

The convalescent plasma, containing number of neutralizing antibodies, reduces the mortality of the severe patients with influenza A, MERS-CoV, and SARS-CoV.¹⁸ The immunotherapeutic potential of convalescent plasma for COVID-19 is also reasonable for the comparable virology and pathology with SARS and MERS.^{40,72} It has been shown that the improvement in 5 critically ill patients with COVID-19 who received 400 mL of convalescent plasma. After the plasma transfusion, the body temperature normalized within 3 days in 4 of 5 patients, PaO₂/FiO₂ increased (from 172–276 pre- to 284–366 post-treatment) within 12 days. 3 patients were weaned from mechanical ventilation within 2 weeks of treatment.⁷³ Duan et al reported that 200mL of convalescent plasma brought improvement of oxygen saturation (median: 93% vs 96%), lymphopenia ($0.65 \times 10^9/L$ vs $0.76 \times 10^9/L$), C-reactive protein (55.98 mg/L vs 18.13 mg/L), and CT scan in 10 patients with severe COVID-19.⁷⁴

Monoclonal antibodies (mAbs) are produced by B cells under the stimulation of specific target antigens.⁷⁵ Most of the antiviral mAbs studied so far are IgG. The viruses or infected cells combine with IgG to form antigen antibody complexes, which bind the FC γ receptors on the surface of killer cells (NK cells, macrophages, etc.). The virus is eliminated by antibody dependent cell-mediated cytotoxicity (ADCC).⁷⁶ Thus, the monoclonal antibodies for the SARS-CoV-2 are effective in elimination of virus. As the relatively high identity of receptor-binding domain (RBD) in SARS-CoV-2 and SARS-CoV, Tian and Yuan found that the SARS-CoV-specific human monoclonal antibody, CR3022, could bind potently with SARS-CoV-2 RBD in vitro.^{77,78} However, potent SARS-CoV-specific neutralizing antibodies of m396 and CR3014 that target the ACE2

binding site failed to bind SARS-CoV-2 spike protein.⁷⁷ Due to its relatively fast approval route and generally favorable safety profiles, mAb therapy should be a promising alternative to the treatment and prevention of emerging viral threats.⁷⁹

Limitations

Although several agents rounding up the immunopathology of COVID-19 have been used clinically, and showed benefits for the patients, such as IFNs, thymosin, glucocorticoids, and IVIG, etc. It is worth mentioning that some drugs which are already granted by Emergency Use Authorization (EUA) have not been involved in our review, including monoclonal antibodies such as Bamlanivimab, Casirivimab, and Imdevimab.⁸⁰

Conclusions

In summary, immune dysregulation, dysfunction and immunopathological injury are crucial to the elimination of virus and the progression of COVID-19. The agents to restore the unbalance of the immune response could be the alternative treatment approach under the scenario of the lacking of effective antiviral drugs.

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