Macrophage Activation and Cytokine Release Syndrome in COVID-19: Current Updates and Analysis of Repurposed and Investigational Anti-Cytokine Drugs

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Introduction
Novel coronavirus disease (COVID-19), originated from the Wuhan city of China has now taken the shape of a pandemic with a total reported case of 37.10 million and deaths of more than 1.07 million across 233 nations [1]. United State of America has a total 7.58 million confirmed cases, followed by India (7.05 million), Brazil (5.05 million), and Russia (1.29 million) as on 10/12/2020 [1]. As per the WHO report, in the USA, Brazil, and Russia, community transmission is confirmed whereas in India, yet community transmission is not reported [1]. Genomic and structural analysis of nCoV-19 has revealed the presence of spike protein, membrane protein, envelope and nucleocapsid with the presence of 10 ORFs (operative response filament) [2]. It was found that the spike protein of the virus interacts with the ACE-2 receptor located at the surface of epithelial cells and alveoli of lungs and enters into the cell via endocytosis [2]. Upon its entry, open reading frames (ORFs), nsp 1-16 and other polyproteins carryout RNA and genomic transcription and further get modified by interacting with endoplasmic reticulum and Golgi bodies. Recent findings have also confirmed the presence of nCoV-19 in the intestine, liver, and in the heart [3, 4]. Common symptoms for COVID-19 infection are cough, fever and pneumonia leading to acute respiratory distress syndrome (ARDS) [5]. In COVID-19 infection, innate immunity plays a vital role in viral clearance. Serological analysis of COVID-19 infected patients has shown increased level of pro-inflammatory cytokines,
which at first instance seems favorable [6]. However, in most of the critical cases, a very high level of pro-inflammatory cytokines have been reported that cause cytokine storm, which is undoubtedly a condition to worry [7]. Therefore, in this manuscript, we have critically analyzed the viral clearance and hyperinflammatory role of cytokines and therapeutic promises of anti-cytokine drugs. Further, we have discussed various mechanism of hyper inflammation that triggers ARDs towards multi-organ damage. At the end we have reviewed some of the repurposed and investigational anti-cytokine therapies in terms of their safety and efficacy. A report published in Lancet, have clarified the presence of tumor necrosis factor alpha (TNF-α), interleukin (IL) IL-6, IL-10, granulocyte colony-stimulating factor (G-CSF), granulocyte-macrophage colony-stimulating factor (GM-CSF), MC1, macrophage inflammatory protein 1 alpha (MIP1-α) and IP10 along with IL-17 during COVID-19 infection [5]. According to Pedersen et al., 2020, production of inflammatory cytokines is an adaptive response towards viral clearance but it was found that not all cytokines participate in viral clearance [8]. IL-12, IL-15, IL-21 and interferons (INFα, β and γ) are specific for clearance whereas an increased level of TNF-α, IL-6, IL-17 G-CSF and GM-CSF leads to hyperinflammation and cytokine storm as shown in Fig. 1, [9].

Macrophage Activation, NLRP3 Inflammasome-NF-kB mediated Hyperinflammation and Multi-organ Toxicity

As nCoV-19 enters into the pneumocytes, some viral particles are engulfed by the macrophage, which serves as an initial line of defense followed by the establishment of antigen-presenting cells (APC) via major histocompatibility complex-2 (MHC-2) of T-cells [9]. This event is followed by the co-stimulation and activation of T-helper cells (CD4+ and CD8+ cells) which in turn produces IL-4 and 5 and causes activation of B-cells [9]. Antigen-presenting cells (APCs), co-stimulation, and produced cytokines also activate natural killer (NK) cells [10]. Therefore, antibodies produced by B cells, granulosomes from NK cells, IL-12, 15, 21 and INF-α, β and γ effectively participate in viral clearance [11, 12]. However, another aspect (hyper inflammation) is mediated by macrophage activation, PMN cells, CD4+, CD8+ cells and other signaling pathways that produce inflammatory cytokines that lead to cytokine storm and CRS (cytokine release syndrome) [13]. IL-17, produced via macrophage and polymorphonuclear neutrophils (PMNs) causes recruitment of monocytes and neutrophil leading to production and activation of downstream chemokines and cytokines such as IL-6, IL-8, and TNF-α and MCP-1 [14]. Thus, there are substantial evidence available that shows the overlapping function of immunopathology in COVID-19 infection and therefore the choice of anti-cytokine therapy must be made with caution.

Apart from the immunomodulatory effect of nCoV-2, ORFs of viral particles have been identified that interact with Toll-like receptor (TLRs) 3, 7, 8 and 9 through retinoic-acid inducible gene (RIG-1), cytosolic receptor melanoma differentiation-associated gene 5 (MDA5) and nucleotidyl transferase cyclic GMP-AMP synthase (cGAS) [3, 6]. This interaction recruits MAVS (mitochondrial antiviral signaling protein) and MyD88 (myeloid differentiation primary response protein) that independently activate c-jun N-terminal kinase (JNK) and nuclear factor kappa-light-chain-enhancer of activated B cells (NF-kB), respectively [15]. Activated JNK further activates p38 which causes apoptosis whereas activated NF-kB causes transcription of pro-inflammatory cytokines such as TNF-α and ILs [16, 17]. Siu et al. in 2020 showed the formation of NOD-, LRR- and pyrin domain-containing protein 3 (NLRP-3) inflammasome complex and increase in the expression of calcium channels on the pneumocytic surface [18]. NLRP3 inflammasome complex causes activation of pro-IL-1β whereas calcium channels increase the influx of calcium into the cells that further induces severe oxidative stress, mitochondrial dysfunction and produces inflammatory cytokines [19]. These cytokines, specifically, TNF-α and IL-6 are released into the systemic circulation, increases vascular permeability in the lungs and damage vital organs [20]. Considerable evidence of various organ damage such as heart, liver and kidney in COVID-19 have been reported in many case studies, supported by biopsy report and serological findings [21–23].

Current Available Cytokine-based Interventions

Type-I interferon

SARS-CoV and MERS CoV infections led to a discovery that IFN-α and IFN-β can be a potential drug target for SARS-CoV-2 infection treatment [24]. Zhou et al., 2020 studied 77 patients with confirmed COVID-19 and were given either nebulized IFN-α2b (5 mU b.i.d.), oral Umifenovir (200 mg t.i.d.), or a combination of both [25]. They concluded that IFN-α2b has significant action in reducing the elevated inflammatory markers [25]. Another study by Wang et al., 2020 evaluated IFN-α1b in the form of nasal drops as a prophylactic measure for the healthcare staff working within the hospital [26]. A total of 2944 volunteers were included which were divided into low-risk and high-risk groups according to the exposure to coronavirus [26]. Outcome of the study showed a significant protective effect of nasal drop and showed no serious adverse effect [26].

Interleukin-7

Cytokine storm with lymphopenia and lymphocyte exhaustion are the hallmark of COVID-19 and considered to be responsible for the accumulation of viruses and reduced clearance of infected cells [11]. IL-7 is a major cytokine that is responsible for the reversal of T-cell exhaustion and promotes lymphocyte expansion, thus it becomes alluring that IL-7 may have a major role in the restoration of the immune system [27]. IL-7 also help in the replenishment of the circulation pool (CD4+ & CD8+) of T cells by inducing potent proliferation of naive and memory T cells and also exerts anti-apoptotic activity [27]. Recombinant IL-7 (rIL-7) rIL-7 when used to treat T-cell exhaustion following septic shock in HIV patient, it was found to be therapeutically beneficial and it restored CD4+ T cell with no evidence of hyperinflammatory response [28]. Thus, use of IL-7 appears to be future therapeutics for the treatment and management of COVID-19 but as of now, there are no registered trials for evaluation of this strategy.
**Fig. 1** Showing immunopathology of SARS-CoV-2. Upper panel showing the mechanism of viral clearance. Engulfed particles are presented at MHC-II and also identified by PMN cells leading to the formation of APC (antigen-presenting cells) which further interact with TH cells (T helper cells) and causes co-stimulation. Co-stimulated TH cells in turn produce IL-4 and 5 that further activate B cells to produce antibodies. Activated TH cells also produce NK cells (natural killer cells) and together participate in the viral clearance. Middle panel shows the mechanism of hyper inflammation and cytokine storm. Viral particle causes recruitment of MVAS and MyD88 that causes increased activity of MAPK kinase and nuclear translocation of NF-kB leading to production p53, IL-6, TNF-α and pro IL-1β. Viral particles also cause the formation of NLRP3 inflammasome that activate pro-IL-1β to IL-1β. These events cumulatively cause cytokine storm and severe ARDs (acute respiratory distress syndrome) and in due course of time, these pro-inflammatory cytokines are released into the systemic circulation and cause multiorgan damage, as depicted in the lower panel.

### Current Available Anti-cytokine Interventions

**Interleukin-6 inhibition**

IL-6 is a major mediator of cytokine storm and its serum levels have been closely correlated with the severity of ARDs and the outcome with SARS-CoV-2 viral load [20]. In a meta-analysis, 2.9 times higher serum levels of IL-6 have been reported in patients who had complicated COVID-19 as compared to patients with non-complicated one [29]. Therefore, inhibition of IL-6 or its receptors antagonism has been found effective in the treatment of cytokine storm associated with COVID-19 complications [30]. IL-6 inhibitors include Siltuximab, Clazakizumab, Sirukumab while its receptor inhibitors
include Tocilizumab and Sarilumab [31]. These are approved drugs in different clinical conditions [31]. A study by Luo et al., 2020 reported increased level of IL-6 in patients with aggravated or fatal outcomes [32]. This study included 21 patients from China where 8 patients received methylprednisolone, 3 patients died while 2 experienced worsenings of the disease, 1 patient improved while 9 of them achieved clinical stability [32]. Xu et. al., 2020 reported the therapeutic effect of Tocilizumab on 21 severe or critical patients of COVID-19 [33]. This study didn’t involve any control group and outcome showed a decrease in oxygen requirements (75 %), improvement in CT scan anomalies (90.5 %) and overall clinical improvement (100 %) with no deaths or adverse effects [33]. A study conducted in Italy reported clinical improvement in 33 % of patients with Siltuximab at doses ranging from 700 to 1200 mg [34]. Rounier et al., 2020 reported the outcome of Tocilizumab in 30 patients with severe pneumonia and Tocilizumab significantly reduced the requirement of ventilator and risk of ICU admission [35]. Thus, looking into the benefit of IL-6 inhibitor in COVID-19, recently, National Health Commission of China’s COVID-19 diagnosis and treatment program (7th edition) included Tocilizumab in patients with extensive bilateral lung lesions opacity whereas the Infectious Diseases Society of America (IDSA), recently, published a guideline to use Tocilizumab in context with clinical trials only [36].

Interleukin-1 family: IL-1 and IL-18 blockade

A set of studies on patients with COVID-19 induced pneumonia lead to the finding that IL-1β and its natural antagonist (i.e., IL-1Ra) is present in the peripheral blood and bronchoalveolar lavage fluid (BALF) [37, 38]. Further, it was found that NLRP3, an inflammasome, which has been most extensively studied in MERS-CoV and SARS-CoV infection, is found to be responsible for activation and secretion of IL-1β [39]. A recombinant IL-1Ra, Anakinra, has proved its efficacy in treating Hemophagocytic lymphohistiocytosis and is currently used in the pharmacotherapy of IL-1β induced auto-inflammatory disease [40, 41]. Anakinra is being evaluated for its efficacy for COVID-19 in a clinical trial (NCT03332225). Further, a monoclonal antibody, Canakinumab, is being investigated for its action on IL-1β in a single-arm observational study (NCT04348448) [31]. Another upregulated cytokine, IL-18, has been observed in BALF of COVID-19 patients and mainly produced by macrophages upon the action of inflammasome-activated caspase-1 [39]. Recombinant IL-18 binding protein, Takedinig Alfa, has shown its efficacy for the abovementioned conditions but as of now, there are no registered clinical trials to analyze its efficacy [42].

Tumor Necrosis Factor-α inhibition

Blood and tissue of COVID-19 patients have shown evidence of higher levels of TNF-α in patients with severe ARDS [43]. TNF-α inhibitors are well-evaluated for their efficacy in COVID-19 [44]. Neutralization of TNF-α in rat models showed protection against SARSCoV infection and a rapid decrease in IL-6 and IL-1 is observed using anti-TNF in rheumatoid arthritis patients [45, 46]. TNF-α inhibitor is also responsible for the reduction of adhesion molecules and vascular endothelial growth factor, affecting capillary leak [47]. Therefore, TNF-α inhibitors have the potential to be a future therapeutic against COVID-19 infection.

Kinase inhibitors

To take care of COVID-19 situation, various anti-inflammatory class of drugs are under trial and among them kinase inhibitors have emerged as one of the potential therapeutic modality to manage and treat COVID-19 infection [48]. These kinase inhibitors have shown ‘one-drug-multiple target’ properties and act on multiple proteins of the COVID-19 virus and hence, inhibit the viral entry, replication as well as mitigate the cascade of the cytokine storm [48]. As most of the cytokine inhibitors are approved for other indications, hence possess established safety and pharmacokinetic profile and therefore appears to be potential drug candidate for repurposing. Thus, looking into these facts, kinase inhibitors, if found potent in mitigating COVID-19 induced complications would significantly minimize the time as well as the cost of drug development and can be accessible to the patients in a short span of time, as compared to an entirely new class of drugs [48]. Recently, Bouhaddou et al., 2020 published an extensive study showing the significant involvement of kinases in the pathogenesis of COVID-19 and thus, opened an avenue to target these kinases against COVID-19 infection [49]. Bouhaddou et al., 2020 have shown the involvement of CK2 (casein kinase II), mTOR (mammalian target of rapamycin), ERK (extracellular signal-regulated kinase), p38 MAPK (mitogen-activated protein kinase), CDKs (cyclin-dependent kinase), EGFR (epidermal growth factor receptor) and ROCK (Rho-associated protein kinases) in the pathogenesis and thus, their inhibitor appears to be the potent anti-COVID-19 therapeutics [49]. As of now many of the kinase inhibitors are under the preclinical stage and many others are under clinical trials. Siltimaserib, a CK2 inhibitor is under Phase 2 trial, Giltekinib (AXL inhibitor, FDA approved), Bencinentib (AXL inhibitor, FDA approved), (AXL inhibitor, clinical trial), ARY-797 and Ralimetinib (p38 MAPKs inhibitor, Phase 2/3 and Phase 2), MAPK13-IN-1 and SB203580 (p38 MAPKs inhibitor, under preclinical investigation) [50].

Janus kinase (JAK) inhibitors

JAK-STAT pathway has been well-documented in the pathogenesis of COVID-19. It plays a critical role in viral entry, viral replication and in the production of various pro-inflammatory cytokines, such as ILs, TNF-α, GM-CSF and interferon-gamma [51]. Thus, JAK inhibitors, mainly JAK1 and JAK2 have been explored for the potential antiviral and anti-inflammatory effect in COVID-19 infection [52]. Among some of the established JAK inhibitors such as Baricitinib, Lestaurtinib, Ruxolitinib, Midostaurin, Lestaurtinib, Ruxolitinib, Fedratinib, and Tofacitinib, Baricitinib (an approved drug for rheumatoid arthritis) has shown potent protection against COVID-19 infection [48, 53–55]. Interestingly, Baricitinib when used at the same dose which was approved for rheumatoid arthritis, showed inhibition of AAK1 (P2-associated protein kinase-1) and GAK (cyclin G-associated kinase) which resulted into inhibition of viral entry via ACE-2 receptors and showed potent anti-inflammatory effect leading to reduction in cytokine storm [55]. Thus, considering the therapeutic promise of Baricitinib, a Phase 2 clinical trial among 800 patients is going on where 2 mg of Baricitinib for 10 days is being assessed for clinical safety and efficacy among COVID-19 infected patients (NCT04321993) [56]. Another Phase 2/3 clinical trial is going on where (NCT04320277) Baricitinib at the dose of 4 mg/day in combination with Lopinavir or Ritonavir at the dose of
250 mg/bid for 14 days is being evaluated for clinical efficacy [56]. Recently, Adaptive COVID-19 Treatment Trial 2 (ACTT-2) (Phase 3), where a combination of Baricitinib and Remdesivir were used, showed a significant reduction in recovery time among COVID-19 hospitalized patients [NCT04401579]. Thus, it can be concluded that JAK-inhibitors could be a potential drug to treat COVID-19 infection via its multifactorial mechanism of action.

**EGFR and CDKs inhibitors**

As it is well documented that pulmonary fibrosis is the clinical manifestation of COVID-19 infection and recently published evidence has shown significant pulmonary fibrosis among post-recovery patients [57], EGFR signaling is considered as the major contributor to fibrosis via modulation of the TGF-β signaling pathway [58]. Further, TGF-β is among the commonly reported pro-inflammatory cytokines among COVID-19 infected patients that aggravate lungs injury and ARDs. Sorafenib, a potent kinase inhibitor, as well as Osimertinib, an EGFR inhibitor, have shown significant anti-inflammatory and anti-fibrotic effect hence became a future therapeutics against COVID-19 infection [59]. Apart from EGFR inhibitor, CDK inhibitor i.e, Atembiclib (an approved drug for advanced breast cancers) showed a significant anti-COVID-19 property and enlisted among the 24 FDA-approved drugs that possess potent inhibitory activity against the COVID-19 virus [48, 60].

**MEK inhibitors**

MEK/ERK has been reported to play a significant role in the pathogenesis of COVID-19 infection [61]. Viral spike protein and nucleocapsid proteins have been reported to stimulate the phosphorylation of ERK/MEK that triggers the release of various pro-inflammatory cytokines leading to ARDs and cytokine storm [62]. ATR-002 is a small molecule and inhibitor of Raf/MEK/ERK pathway [63]. Apart from the preclinical evidences, the outcome of the Phase I clinical trial showed the safety and clinical efficacy of ATR-002 in terms of reduced virulence and cytokine storm (NCT04385420) [63]. Further, In Vitro study using ACE2-A549 has shown p38 MAPKs activation in response to COVID-19 infection and blockage of p38 MAPK pathway by its inhibitor SB203580 showed a significant reduction in the level of IL-6, TNF-α and other inflammatory cytokines [49]. Apart from SB203580, ARRY-797 and Ralimetinib (p38 MAPKs inhibitor, Phase 2/3 and Phase 2), MAPK13-IN-1 (p38 MAPKs inhibitor, preclinical) are also potential MAPK inhibitors and could be a future therapeutics against COVID-19 infection [49].

**Analysis of Repurposed and Investigational Anti-cytokine Drugs**

As it seems very persuasive that cytokine storm is a major factor in worsening the clinical outcome of COVID-19 infection and therefore use of anti-cytokine therapy seems to be a better option. However, it must be kept in mind that there still exists a fine line of difference between viral clearance mechanism and hyper inflammation and we need to specifically stimulate the viral clearance pathway and block the hyperinflammatory pathway [64]. Looking into the inflammatory role of IL-6 and TNF-α, anti-IL-6 (Tocilizumab, Sarilumab) and anti-TNF-α antibodies (Infliximab, Etanercept, Golimumab and Adalimumab) have been used and outcomes were promising [8, 65]. At present these drugs are under clinical trial (phase I and II). JAK-STAT pathways is yet another important signaling pathway that regulates immune and inflammatory response [66]. IL-6 has been reported to interact with JAK-STAT and activate this cascade [27]. Thus, apart from using, anti-IL-6 antibodies, JAK inhibitors can also be a novel therapeutic approach. However, use of JAK1 and 3 inhibitors have been reported to interact and inhibit IL-15, IL-21 and INFs, which is essential for viral clearance and inhibition of these cytokines could be fatal [9]. However, the use of the JAK2 inhibitor has been reported to inhibit IL-17 (responsible for cytokine activation), IL-6 and GM-CSF which are responsible for cytokine storm [67]. This seems to be a favorable approach. Apart from the aforementioned drugs, several other anti-inflammatory drugs such as NSAIDs, Sulfasalazine, Mesalazine and COX-2 inhibitors are being investigated [3, 68] but we strongly suggest to take care while selecting these anti-cytokine/anti-inflammatory drugs, as slightest of deviation can be fatal for patients.

**Conclusion and Future Direction**

Looking into the pathological role of cytokine storm and previous background of anti-cytokine activity of anti-IL-6 or TNFα antibody, we speculate that the use of these drugs can be beneficial but use must be on a case to case basis. We also need to think of two aspects regarding the use of these drugs (1) when anti-cytokine therapy should be given? and (2), how safe these drugs are?. Based on the published reports, it can be said that early use of anti-cytokine therapy will be a better option, as compared to steroidal therapy which is meant for immunosuppression [69]. Use of these drugs should be avoided in case, patients are in intensive care. Concerning the safety, use of high dose anti-inflammatory drugs, in general causes tissue damage and potent anti-inflammatory drugs increase the risk of supra infection (bacterial and fungus infection). However, till now, no such case of supra-infection have reported. Therefore, in nutshell, anti-cytokine drugs have the potential to be a choice of drugs for the management and treatment of COVID-19 associated ARDs and cytokine storm but large sample size study will only decide their forthcoming future. Until now, for a healthy individual, social distancing along with an active lifestyle and for critically ill patients, plasma therapy appears to be a safer alternative, unless vaccine is introduced.

**Conflict of Interest**

The authors declare that they have no conflict of interest.

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