TLRs in COVID-19: How they drive immunopathology and the rationale for modulation

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Abstract
COVID-19 is both a viral illness and a disease of immunopathology. Proximal events within the innate immune system drive the balance between deleterious inflammation and viral clearance. We hypothesize that a divergence between the generation of excessive inflammation through over activation of the TLR associated myeloid differentiation primary response (MyD88) pathway relative to the TIR-domain-containing adaptor-inducing IFN-β (TRIF) pathway plays a key role in COVID-19 severity. Both viral elements and damage associated host molecules act as TLR ligands in this process. In this review, we detail the mechanism for this imbalance in COVID-19 based on available evidence, and we discuss how modulation of critical elements may be important in reducing severity of disease.

Keywords
TLR, COVID-19, innate immunity, interferon, CD14

The innate immune system's antiviral response
Upon infection with SARS-CoV-2, the virus that causes COVID-19, an innate immune response is activated through the phylogenetically preserved TLR family. TLRs are germline encoded PRRs that are believed to be important for both the protective and maladaptive response to lower respiratory viral infection. Each TLR recognizes ligands comprised of both PAMPs elaborated by the virus and damage associated molecular patterns (DAMPs) released by host injured and dying cells. The leukocytes of the innate immune system, including neutrophils, monocytes, and NK cells, depend on TLR signaling to appropriately localize to the site of infection and target virus and virally infected host cells. This initial response helps control viral infection but also causes cell damage and release of DAMPs, further TLR activation, and more inflammation. This cycle is thought to drive organ injury. TLR binding is necessary for the maturation of APCs through production of pro-inflammatory cytokines (e.g. TNF-α), chemokines, chemokine receptors, and co-stimulatory molecules. These early innate immune responses initiate and direct the subsequent Ag-specific adaptive immune responses necessary for viral clearance. This leads to a race between ongoing damage from direct viral infection and the innate hyperinflammatory response versus the protective innate and adaptive antiviral responses (Figure 1).

In viral infection, it has been demonstrated that the cell surface receptors TLR1/2, TLR2/6, and TLR4 recognize structural components including proteins and glycolipids. TLR3, TLR7, TLR8, and TLR9 are expressed on endosomes and recognize nucleic acids. Each TLR has an extra-cellular leucine-rich repeat domain and an intracellular Toll/IL-1 receptor (TIR) domain. Upon TLR binding the TIR domain recruits adaptor proteins which leads to an MyD88- or TRIF-dependent signaling cascade. The MyD88-dependent pathway generates pro-inflammatory cytokine (IL-1, IL-6, TNF-α, etc.) production via transcription factor NF-κB. Alternatively, the TRIF-dependent pathway leads to activation of the type 1 and type III IFN critical for viral defense in addition to inflammatory cytokines. Depending on the TLR, cell type and subcellular...
compartment, downstream signaling through either MyD88 or TRIF can generate both pro-inflammatory cytokines and a type I/III IFN response. The heterodimers TLR2/6 and TLR2/1 are MyD88-dependent as cell surface receptors but can traffic to endosomes and generate a type I IFN response. TLR3 recognizes dsRNA and signals solely through the TRIF pathway to generate a type I/III IFN response. TLR4 uses both MyD88- and TRIF-dependent pathways to generate a variable response. The prototypical TLR4 ligand is LPS but other ligands including glycolipids, viral structural proteins, and DAMPs such as HMGB1 have been identified. TLR7 and TLR8 recognize ssRNA and TLR9 recognizes unmethylated CpG DNA. TLR7 and TLR9 are MyD88-dependent but effectively generate a type I/III IFN response. Other PRRs including the cytosolic RNA sensing RIG-I-like receptors RIG-I and MDA5 also participate in viral recognition and response through TLR-independent mechanisms. Therefore, upon infection with SARS-CoV-2, viral particles bind different TLRs and other PRRs and activate the innate immune system. This response consists of the generation of inflammation through transcription factor NF-kB and an antiviral response through the TRIF pathway.

The antiviral properties of TLR induced type I and type III IFN are critical for coordinated viral control and clearance. Type I IFN consist predominantly of IFN-α and IFN-β and these cytokines signal through a common receptor IFNAR1/2. IFNAR1/2 binding leads to signal transduction via the JAK/STAT pathway and eventual formation of the transcription factor ISGF3 and expression of IFN-stimulated genes (ISGs). Type III IFN consist of the cytokines IFN-λ1/2/3 which are structurally unique from IFN-α and IFN-β but exhibit similar functions; the main difference is that type III IFN receptors primarily localize to epithelial surfaces, for example airway epithelial cells. When type III IFN bind their receptor, similarly, signal transduction is initiated through the JAK/STAT pathway with eventual formation of ISGs. Protein kinase R (PKR) and 2′-5′-oligoadenylate are two well-characterized ISGs that act on translational machinery and nucleic acids directly to impair viral replication. MERS-CoV, another coronavirus with high sequence homology to SARS-CoV-2, produces proteins that antagonized PKR so as to subvert the host antiviral response. Within the innate immune system IFN production activates DCs, NK cells, and macrophages to directly clear and kill virally infected cells. IFN

Figure 1. (a) Schematic detailing the innate immune response in COVID-19 beginning with viral infection of type II pneumocytes. Viral elements (PAMPs) and DAMPs are generated and act as ligands for TLR activation. Several cell surface and endosomal TLRs are activated. Through MyD88-dependent and TRIF-dependent pathways, an NF-kB derived inflammatory response and an IFN response are generated. (b) NF-kB derived inflammation generates DAMPs and initiates a cycle of innate immune activation. This inflammation may drive organ injury. The IFN response is critical for viral clearance. As both pathways are active, a race between viral clearance and host injury ensue. In severe disease, the balance is shifted toward injurious NF-kB derived inflammation. DAMP: damage associated molecular pattern.
also play an immunomodulator role on APCs and lymphocytes to generate an effective adaptive immune response. In vitro, SARS-CoV-2 is sensitive to IFN-α pre-treatment; specifically, when Vero and Calu3 cells are pre-treated with IFN-α, SARS-CoV-2 infection is unable to halt phosphorylation of STAT and downstream expression of ISG proteins resulting in resolution of SARS-CoV-2 infection. However, there is limited data about the timing of TLR mediated type I IFN induction in COVID-19.

Concurrently to this TLR-mediated IFN response, TLR activation generates a robust inflammatory response through the MyD88-dependent pathway and transcription factor NF-κB. This pro-inflammatory response leads to an increase in cytokines and chemokines and an influx of activated immune cells (neutrophils, monocytes, NK cells) to the site of infection. Patients with severe COVID-19 compared to uninfected patients have higher serum levels of pro-inflammatory cytokines and chemokines including IL-2, IL-6, IL-1β, IL-8, IL-17, G-CSF, GM-CSF, IP10, MCP1, MIP1α, and TNF-α. This pro-inflammatory state and direct infection by virus lead to host cell death and organ injury. Damaged host cells release DAMPS like HMGB1 and oxPAPC that are capable of amplifying inflammation by TLR binding particularly through the MyD88-NF-κB pathway. Pro-inflammatory states are typically accompanied by an increase in anti-inflammatory cytokine production, serving as a negative feedback loop. IL-10 is a negative regulator of the immune response producing a negative feedback loop.

In COVID-19, and it is higher in critically ill patients. COVID-19, and it is higher in critically ill patients. The notion of a TLR mediated dys-regulated immune response was previously investigated in SARS. In a mouse model of SARS-CoV-1 infection, it was found that MyD88 knockout mice and TRIF knockout mice had similar mass loss and mortality. In this same study, the TRIF knockout mice had a more robust cytokine response and a higher viral load in type II pneumocytes. Additionally, it was found that TLR3 knockout mice, TLR4 knockout mice and TRAM (TLR4-TRAM-TRIF pathway) knockout mice all lost mass findings have been observed across experimental methods including in vitro tissue culture, ex vivo infection of primary cells, and in vivo samples derived from patients and animal studies. However, in contrast to these findings, Broggi et al. found that in COVID-19 disease severity correlated with increased IFN-λ expression within the lungs. Further, in an animal model, they found that exposure of murine lungs to a synthetic RNA virus led to lung DC IFN-λ production. This then caused epithelial barrier damage and an increased susceptibility to bacterial co-infection. One proposed explanation for these discordant findings is that while local type III IFN production in the lungs may contribute to immunopathology, systemic type I and type III IFN production may be protective. IFN is being actively studied as a therapeutic in COVID-19. Intra-nasal IFN-α prophylaxis in addition to PPE reduced the incidence of SARS-CoV-2 infection in at risk personnel. In severe COVID-19, in a small randomized controlled trial, systemic IFN-β therapy improved discharge rate and reduced mortality. In another randomized control trial, IFN-β in combination with lopinavir-ritonavir and ribavirin compared to lopinavir-ritonavir alone reduced time to negative nasopharyngeal SARS-CoV-2 PCR result. In a non-randomized retrospective cohort study, IFN-α or without the antiviral arbidol improved time to viral clearance and reduced systemic inflammation. Interestingly, in COVID-19 patients, IFN stimulation appears preserved when exogenously stimulated by other agonists; this raises the question of how upstream proximal factors in TLR activation may impact the adaptive immune response in COVID-19.

One possibility is that the virus itself may possess an evasion mechanism to protect against viral dsRNA and ssRNA recognition by TLRs 3 and 7 as was the case with SARS-CoV-1. Han et al. report that the SARS-CoV-2 accessory protein ORF9b impairs induction of types I and III IFN in part by targeting components of the TLR3-TRIF endosomal RNA sensing pathway in vitro. Yin et al. report in vitro data describing that the PRRs MDA5 and LGP2, as opposed to TLR3, primarily recognize SARS-CoV-2 RNA and regulate a delayed IFN response in airway epithelial cells. TLR3 and 7 are critical activators of the type III IFN antiviral response, and these data suggest SARS-CoV-2 evasion. Others suggest that DEAD-box RNA helicases in humans may induce a TLR independent anti-viral IFN response to SARS-CoV-2 though evidence is lacking.

The immune response to SARS-CoV-2

In COVID-19, severe disease correlates with an immune signature characterized by an early cytokine and chemokine rich inflammatory response likely through NF-κB and very low level of antiviral IFN gene activation. This has been described in patients, in whom a low ISG signature in peripheral blood correlated with disease severity; additionally, in patients with severe disease an increased SARS-CoV-2 viral load was observed suggesting delayed viral clearance. In this same study, elevated NF-κB pathway expression of IL-6 and TNF-α was also observed. Similar
TLRs in COVID-19

TLR4

TLR4 is positioned to respond to both exogenous PAMPs and DAMPs released as part of tissue injury. TLR4 cooperates with MD-2 and CD14 to bind extra-cellular ligands. TLR4 signals both through the MyD88-dependent and TRIF-dependent pathways. LPS, a component of most Gram-negative bacterial cell envelopes, is its classic ligand. TLR4 activation has been implicated in the development of a pro-inflammatory state as in Gram-negative bacterial sepsis. This response is critical to host defense in Gram-negative bacterial infection. For example, mice strains with TLR4 genotypes that lead to LPS hypo-responsiveness have increased susceptibility to Gram-negative infections. However, excessive LPS directed TLR4 stimulation can lead to damaging inflammation. In fact, the administration of lipid A (a component of LPS), is extremely toxic at high doses and leads to a hyper-inflammatory response and fulminant sepsis-like syndrome in animals. This response is thought to be driven primarily through MyD88 activation as opposed to TRIF signaling.

In COVID-19, it is proposed that the viral structural proteins and glycolipids are PAMPs that bind TLR4 initiating an innate immune response. Computer-based modelling has found that the S protein of SARS-CoV-2 is predicted to bind to TLR4. Further, emerging data shows that purified SARS-CoV-2 S protein binds TLR4 with strong affinity in vitro, and that purified S protein induces a TLR4-dependent IL-1β response akin to the response observed with LPS challenge. The balance in downstream activation of MyD88 versus TRIF-dependent pathways is unknown. However, higher MyD88 activation relative to TRIF activation may contribute to excess NF-κB activation and inflammatory cytokine production as TLR4 has been heavily implicated in the injurious inflammation and lung injury seen SARS-CoV-1 and MERS. TLR4 modulation or blockage has been shown to reduce lung injury in animal models by reducing NF-κB pathway activation. Further work is needed to determine the extent by which TLR4 binding of viral PAMPs including the SARS-CoV-2 S protein drives lung injury and viral sepsis in COVID-19.

In addition to recognition of viral proteins, several DAMPS also act as ligands for TLR4. Viral infection and subsequent inflammation lead to the generation of DAMPs. For example, HSP60 and HSP70, released from virally infected and stressed cells are TLR4 agonists. In addition, fibrinogen, an acute phase reactant, binds and activates TLR4. HMGB1 is a DAMP that binds TLR4 and has been studied in the context of influenza-associated ALI. OxPAPC, a heterogeneous group of lipids (oxidized phosphorylcholine derivatives) released from dying virally infected cells, is another DAMP that has been shown to promote inflammatory responses. TLR4 binding of oxPAPC in cooperation with CD14, generates inflammatory cytokines at levels akin to LPS binding. Both the MyD88 and TRIF pathways are activated by DAMPS but there may be a bias toward MyD88 signaling and inflammatory cytokine production. In several viral pneumonia models, TLR4 blockade led to a reduction in inflammation, DAMP accumulation and ALI suggesting that TLR4 activation preferentially generates an injurious pro-inflammatory state.
TLR4 binding may impact the characteristics and efficacy of an adaptive immune response. The TLR4 pathway is capable of elaborating type I/III IFN which are necessary for a virus specific adaptive immune response. Past work has shown that the dose of LPS signaling through TLR4 differentially regulates a Th1 versus Th2 response. However if TLR4 binding leads to excess cytokine production, for example il-6 as is seen in COVID-19, cytotoxic CD8+ T cells and cell mediated immunity may be impaired. To date, the effect of IL-6 blockade on clinical outcomes in COVID-19 has been mixed.

**TLR7**

Functional TLR7 genetic variants have been linked to COVID-19 severity in multiple studies, strongly suggesting a key role for TLR7 in COVID-19 pathogenesis. TLR7 binds ssRNA and imidazoquinolones. While TLR7 signals through the MyD88-dependent pathway, downstream signaling leads to both NF-κB mediated inflammation and a type I/III IFN response; the balance between NF-κB mediated inflammation and the IFN response is critical. Whole exome sequencing in families with multiple cases of COVID-19 identified rare loss of function variants in TLR7. Further, *in vitro* studies in these patients revealed that type I/III IFN responses to imiquimod, a strong TLR7 agonist, were greatly down-regulated in peripheral blood mononuclear cells. These data suggest aberrant type I/III IFN responses due to a functional genetic mutation are specifically tied to severe COVID-19. Given that the gene for TLR7 is found on the X-chromosome it has also been proposed that TLR7 might play a role in sex-based differences in COVID-19 susceptibility and severity. Men are nearly two times as likely to develop respiratory failure or die from COVID-19. Incomplete X-inactivation in immune cells resulting in an increased gene-dosage of TLR7 in women relative to men could lead to a more effective early antiviral response. Common variants in TLR7 have also been identified in association with severe COVID-19 in an exploratory study using multilevel filtering (e.g. frequency, chromosomal, geographic, and functional filters).

Imiquod, a TLR7 agonist, is currently under investigation as an immune modulator in COVID-19. Imidazoquinolones have well-described antiviral properties through activation of TLR7 and the MyD88-dependent pathway with downstream type I/III IFN generation. Imiquod has been previously studied for its antiviral effects in anal condylomata in HIV. In addition to its antiviral properties TLR7 activation may decrease inflammation. In an asthma model, TLR7 agonist RSQ reduced allergen induced airway inflammation and decrease reactive oxygen species. Interestingly, inhibition of MyD88 reversed RSQ-mediated effects. It has also been studied as an immune modulator and as a vaccine adjuvant in influenza.

Additionally, Rutin, a flavonoid compound, that effectively binds TLR7 (and TLR2/TLR6) has been identified as a potential immune modulator. In addition to binding TLRs, rutin binds and potentially blocks SARS-CoV-2 main protease (Mpro). Mpro is an important enzyme that mediates viral replication and transcription in coronaviruses, and it has been identified as an attractive target in drug development. Rutin may play a role in both immune modulation via TLR agonism and an inhibitor of viral replication via Mpro antagonism. TLR7 is emerging as a proximal element in the immune response to viral infection that when activated may lead to downstream elaboration of a net antiviral response.

**TLR3**

TLR3 has the potential to improve both innate and adaptive immune responses in COVID-19 through type I/III IFN production. TLR3 agonists include dsRNA and poly IC, a synthetic RNA analogue. It is presumed to be critical in the recognition of respiratory RNA viruses like SARS-CoV-2; during viral replication, dsRNA is generated and stimulates endosomal TLR3 in addition to other intracellular receptors (RIG1/MDA5). Unlike all other TLRs, TLR3 is TRIF dependent and MyD88 independent. Stimulation of TLR3 results in recruitment of the TRIF adaptor protein and downstream activation of ISGs. TLR3 does not directly activate the pro-inflammatory NF-κB pathway. Because TLR3 activation favors type I/III IFN production, it is thought to be critical in viral defense. In fact, TLR3 mutations and deficiency have been described as monogenic drivers of other viral illnesses. Patients with TLR3 mutations and deficiency are highly susceptible to pediatric herpes virus encephalitis, and patients with TLR3 deficiency are at higher risk of developing ARDS in Influenza A viral infection. Fortunately, in these influenza ARDS cases, *in vitro* type I IFN therapy, the downstream product of TLR3 activation, reduced the susceptibility of airway epithelial cells and fibroblasts to influenza A viral infection suggesting modulation may be helpful.

Antiviral properties of TLR3 and disease modulation with TLR3 agonists have been described in several respiratory viral infections including SARS-CoV. In a SARS-CoV mouse model, TLR3/TRIF signaling played a critical role in the innate immune response through ISG protein production. Additionally, in aged mice, intra-nasal poly IC protected mice from lethal SARS-CoV. Synthetic TLR3 agonists have been studied in the context of numerous other viral infections in animal models. In fact, Poly IC or modified forms has been protected against the following viral infections in animal models: influenza virus, Herpes simplex virus type 2, western equine encephalitis virus, and others.
yellow fever virus, RSV, and others. While these data suggest a protective role, the use of Poly IC has been largely limited by systemic toxicity. However, scientists have investigated modifications to limit adverse effects, and modified versions have been safely used as vaccine adjuvants. The protective role of synthetic Poly IC is currently being investigated in a phase 1 trial in COVID-19 (NCT: 04672291).

**TLR2 heterodimers**

The TLR1/TLR2 heterodimer binding by SARS-CoV-2 S protein may contribute to the hyper-inflammflammatory state and lung injury seen in COVID-19. TLR2 forms heterodimers with TLR1 and TLR6, respectively, to create functional cell surface receptors. In vitro and in vivo studies show that TLR2 recognizes the SARS-CoV-2 envelope protein resulting in MyD88-dependent inflammation. In silico studies suggest that TLR1, TLR2, and TLR6 effectively bind the viral S protein in SARS-CoV-2. In particular, specific peptides have been identified within the S protein as TLR2 agonists. It is not clear to what extent S protein binding of these heterodimers impacts the immune response in COVID-19. While these heterodimers typically signal via the MyD88 pathways, in certain immune cells TLR2 may be internalized and localize to endosomes facilitating a type I/III IFN response. These specific peptides and TLR ligands within the S protein are being further investigated as SARS-CoV-2 specific and antigenic molecules that may be used in vaccine development.

As noted, TLR1 acts exclusively as a heterodimer with TLR2. In sepsis, TLR1 polymorphisms, in particular TLR1-7202G, may impact patient outcomes including the predisposition to the development of ARDS. Further in trauma related-sepsis, there was a trend towards a higher incidence of ARDS among patients homozygous for TLR1-7202G. Using translation methods, it was shown that a specific TLR1 polymorphism was associated with increased TLR1 expression, TLR1 mediated NF-κB activity, and cytokine production. TLR1 polymorphisms may also impact the regulation of the pro-inflammatory state through their influence on regulatory T cells. This data suggests that functional variants in TLR1 lead to variable cytokine expression and lung injury. The SARS-CoV-2 S protein is predicted to bind TLR1/2; TLR1/2 agonism may contribute to the pathophysiology of cytokine storm and lung injury seen in patients with severe COVID-19.

The TLR immunomodulator PUL-042 is under investigation in COVID-19. PUL-042 is a combination drug comprised of the TLR2/6 ligand Pam2CSK4 and the TLR9 ligand ODN M362. It was developed in order to augment host response to respiratory infections. In fact, in animal models it has been shown to effectively improve survival and reduce pathogen burden in otherwise lethal SARS-CoV and MERS-CoV infections.

**TLR5**

TLR5 is most relevant in viral vaccine development as an adjuvant. TLR5 is activated by bacterial flagellin. It has been studied for immunotherapeutic development in respiratory infections because of its presence on respiratory tract epithelium; for this reason, it has been proposed as a candidate target in COVID-19 vaccine and drug development. Administration of flagellin has been previously studied in influenza virus; in a mouse model, flagellin administration reduced viral replication in an interferon-independent manner. Flagellin administration also enhanced the efficacy of oseltamivir. Interestingly, the protective effects appears to be independent of interferon. Flagellin has been a successful adjuvant in several viral vaccines including IAV flu models although its use has been limited by toxicity. In COVID-19, S protein multi-epitope vaccines have been developed with antigenicity confirmed by the degree of TLR5 docking.

**CD14, a TLR accessory molecule**

As a proximal regulator of multiple TLRs, CD14 modulation may impact inflammation and organ injury in COVID-19. CD14 is a protein that is both membrane-bound on immune cells and soluble. It has a large flexible hydrophobic pocket and several grooves that are accessible for ligand binding. It acts in concert with TLR1/TLR2, TLR2/TLR6, TLR4, and TLR9 by binding a variety of PAMPS and DAMPs. Its role in as a co-receptor for nucleic acids for TLR3 and TLR7 is less certain in humans.

CD14 may facilitate an initial innate immune system response to PAMPS like protein S and other viral structural proteins and glycolipids. However, with disease evolution it also likely recognizes DAMPs in addition to PAMPS leading to a net increase in immunopathology. As noted previously, one such DAMP: oxPAPC, an LPS-like DAMP generated from host cell death facilitates innate immune system hyper-activation via pro-inflammatory signaling pathways.

In COVID-19, using a novel ultra-high throughout proteomic approach, up-regulation of CD14 correlated with severity of COVID-19 infection. Further, in COVID-19, soluble CD14 levels were associated with severity of disease. Additionally, an increase in CD14 + CD16 + monocytes were identified in the peripheral blood of critically ill patients with COVID-19. Presepsin, a soluble N-terminal component of CD14, has been identified as a biomarker associated with disease severity in COVID-19. Because of these findings and previous work in sepsis and ARDS, CD14 inhibition has been identified as a potential means to target aberrant innate immune system activation in COVID-19.

It is possible that CD14 is a proximal modulator of several TLRs and plays a central role in regulating the inflammatory response observed in severe COVID-19.
CD14 blockade may impart a net protective effect through modulation of several TLRs. As noted above, CD14 complexes with TLR1/TLR2, TLR2/TLR6, TLR4, and TLR9 to bind PAMPs and DAMPs. However, CD14 has not been definitively shown to interact with TLR3 and TLR7 in humans. TLR3 and TLR7 seem to be critical for an effective antiviral response through the TRIF-IFN pathway. Therefore, CD14 blockade may not decrease the TLR-TRIF-IFN response. TLR1/TLR2, TLR2/TLR6, TLR4, are predicted to bind PAMPs and DAMPs in COVID-19; this may drive an excessive MyD88-NF-κB-dependent inflammation. Therefore, blockade with an anti-CD14 Ab, may actually lead to reduced inflammation while not interfering with the antiviral response. A phase 2, anti-CD14 Ab clinical trial in COVID-19 is currently underway (NCT04391309).

CD14 has been implicated in several deleterious pro-inflammatory disease states: sepsis and septic shock. Previous work has shown that the CD-14 mAb is safe and that when co-administered, less inflammation is generated by LPS challenge. While sCD14 levels increase with Ab administration, inflammatory cytokine levels decrease. CD14 mAbs were previously studied in a phase 1 severe sepsis clinical trial. While CD14- Abs reduced inflammation, the study was underpowered to detect clinical benefit, and CD14 Abs antibodies are not used routinely in clinical practice.

Other TLR accessory molecules in COVID-19

IRAK-1 is a critical proximal component of the TLR MyD88 pathway. It may functionally bridge different pathways including the JAK/STAT- and TRIF-dependent pathways which may be a critical regulator of the pro-inflammatory versus type I/III IFN response. IRAK-1 has been implicated as a potential gene variant critical to the pathogenesis of COVID-19. IRAK-1 has been previously studied in the context of endotoxin shock and autoimmune disease. IRAK-4 also acts within the MyD88 pathway. IRAK-4 mutation has been implicated in recurrent and life threatening bacterial infections in children. Its expression was up-regulated in a small group of patients with COVID-19.

CD24Fc is a biologic immunomodulator. CD24 acts as a checkpoint molecule that modulates TLR4 signaling in response to DAMPS, as in viral infection. For example, upon oxPAPC binding CD24 is recruited to the intracellular TLR4 module and ameliorates or blocks pro-inflammatory signaling. CD24Fc has been designed to reduce pathologic activation of TLR4. While previously studied in the context of GVHD after stem cell transplant, CD24Fc has recently been shown to reduce the development of ARDS in viral pneumonia. CD24Fc is being studied as a therapeutic in COVID-19. Initial reports from a phase III randomized control clinical trial suggest CD24Fc may reduce disease progression, shorten length of stay, and may blunt the immune response. Specifically, CD24Fc treated patients had reduced activation of CD8+ and NK cells and reduced cytokine and chemokine levels.

Conclusion

Upon infection with SARS-CoV-2, TLRs and their proximal accessory molecules play a critical role in both the protective and maladaptive innate and adaptive immune system responses. We hypothesize that the balance between MyD88-NF-κB-dependent excessive inflammation and TRIF-IFN-dependent antiviral response drives severity of disease in COVID-19. The goal of proximal modulation of the innate immune system should be to shift the balance towards a systemic antiviral TLR-TRIF-IFN response and away from immunopathology related to excessive TLR-MyD88-NF-κB activation. As discussed in detail above, one strategy is to inhibit CD14 which acts as a co-receptor for multiple TLRs and in particular those associated with the generation of excessive inflammation including TLR4 and TLR2.

Further investigation into how each TLR contributes to the net immune response and the pathophysiology of COVID-19 is needed. Future work should also be directed at better understanding how modulation of TLRs and accessory molecules impact each phase of the innate and adaptive immune responses to SARS-CoV-2.

Declaration of conflicting interests

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