Human immune response to SARS-CoV-2: What is known? A scoping review

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Abstract
Monitoring literature on the broad spectrum of the human immune response to SARS-CoV-2 is important to understand the mechanisms and progression of COVID-19. The present study undertakes a scoping review of the literature on human immune response to SARS-CoV-2 to determine the characteristics of innate and adaptive responses, as well as biomarkers and cells that play a role in the development of the infection. We searched papers in MEDLINE/PUBMED and EMBASE databases published since December 1st 2019 to April 9th 2020 from which we selected 56 for this study. We found that the immune response is characterized by high levels of acute phase reactants, neutrophilia, low levels of NKs and eosinophils, lymphopenia, cytokine storm syndrome, exhausted T cells, impaired cytotoxic response, inadequate helper response and production of specific antibodies; concluding that immune dysregulation correlates with disease severity and high mortality.

Introduction
In December 2019, an outbreak of a novel coronavirus pneumonia called COVID-19 was first identified in Wuhan, Hubei province, China1–3. As of April 9th, 2020, there have been 1.436.198 cases and 85.522 deaths reported in at least 178 countries4. The coronavirus called SARS-CoV-2 has an incubation period of 0-14 days with a median of 5.2 days5. The clinical spectrum of the infection can vary from asymptomatic or mild clinical symptoms (81% of the cases), and pneumonia of different degrees of severity (14%) to acute respiratory distress syndrome (ARDS) with a high risk of death (5%)5,6.

The most common symptoms are fever, cough, myalgia and fatigue, and the most severe include dyspnea and lethargy5,7. The reason why some people develop severe illness while others do not remains unclear. It is possible that the severity of the disease may be associated with the ability of the im...
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Methods

Protocol and registration: The protocol was written using the elements established in the PRISMA ScR extension document. The research was reviewed by the team members of the Immunology and Translational Medicine group of the National University of Colombia, registered in Colciencias. The protocol was prospectively registered in the Open Science Framework on 13 April, 2020 https://osf.io/t4vw7/

Eligibility criteria: To be included in the review, papers needed to measure or focus on specific dimensions of the human immune response to SARS-CoV-2. Papers in English involving human research, quantitative, qualitative, and mixed design, qualitative review articles, systematic reviews, letter to editors and case reports published from December 2019 to 9 of April 2020 were included in this review. When letters to the editor referred articles of interest that were not found in the initial search, the new articles were also included.

Information sources: To identify potentially relevant documents, the following bibliographic databases were searched: MEDLINE/PUBMED and EMBASE. The search strategies were drafted by the team member discussion and included MESH, emtree and free language terminology. The final search was exported into Mendeley and duplicates were removed.

Search: The final search strategy for PUBMED/MEDLINE and EMBASE can be found in Appendix.

Selection of sources of evidence: The inclusion criteria of the articles were reviewed by title and abstract. To increase consistency among the reviewers, teams of four researchers were generated to evaluate the results of the excluded articles. Disagreement on study selection and data extraction were resolved by consensus and discussion with other reviewers if needed.

Data charting process: A data-charting form was jointly developed by the team to determine which variables to extract. The data was charted independently by each reviewer, the results were discussed by the entire group and the data charting form was continuously updated in an iterative process. Data from eligible studies were charted using a tool designed for this study. The tool captured the relevant information on key study characteristics and detailed information on all metrics used to describe the human immune response to SARS-CoV-2. The data was extracted from the main publications and supplemental materials.

Data items: Article characteristics were extracted (e.g., authors, origin, objectives, purpose), as well as the data related to immune clinical parameters, innate and adaptive immune response, cytokine storm, T cell responses, B cell responses and antibodies. The final version of the charting form can be found in the Additional file Table S1.

Synthesis of results: The studies were grouped by researched items. The results were presented in a narrative format, tables, and figures.

Results

Selection of sources of evidence: The search yielded 240 articles from PUBMED / MEDLINE and 210 articles from EMBASE. After duplicates were removed, a total of 388 citations were identified and 12 indirect articles of comments or letters to editor were added (figure 1). Based on the title and the abstract, 282 were excluded, with 118 full text articles to be retrieved and assessed for eligibility. Of these, 62 were excluded for the following reasons: 5 were in Chinese, 1 was research in animals, 42 not considered the immune response, 3 were vaccine research, 3 were comments or abstracts of original papers and 7 were about possible treatments. Description of the excluded articles can be found in Additional file, table S2. The remaining 56 articles were included in this review.

Characteristics of sources of evidence: The selected articles were classified by the topics shown in Table 1.

Results of individual sources of evidence: The type of publication, journal and aims of the search are presented in additional file: Characteristics of included studies (supplementary table S3). 1 clinical trial, 2 analytic studies, 14 descriptive studies, 9 case reports, 15 reviews, 4 basic research studies and 10 commentaries were included.

Results: For the purpose of this article and due to heterogeneity among articles, the stage of clinical compromise was classified as Severe Patient (SP) if patients were admitted to ICU, progressed to ARDS, died or were referred as severe. If that was not the case, they were considered as Non-Severe Patients (N-SP).

Discussion

Clinical Parameters associated to Immune Response

Usually upon hospital admission in the context of COVID-19, infected patients had abnormal laboratory results. In the case of complete blood count (CBC), the majority of articles that reported lymphocytes and eosinophils counts showed numbers below normal values, and basophils were either normal or decreased. On the other hand, total leukocytes, neutrophils, and monocytes counts displayed a tendency towards normal values. (Figure 2A).

Among cited references, levels of change in CBC varied according to severity of the disease. The majority of references that compared CBC between SP and N-SP reported increased
values of leukocytes and neutrophils in SP\textsuperscript{1,5,7–9,12,18,24,25,33,34,36,41}. Moreover, concurrent with clinical worsening, more than half of the references, reported decreased levels of monocytes, lymphocytes, eosinophils, and basophils in SP\textsuperscript{5,7,9,10,12,18,24,25,33–36,41} (Figure 2B). Additionally, it has been proposed that higher neutrophil-lymphocyte-ratio and higher platelet-lymphocyte ratio are associated with poor prognosis\textsuperscript{9,17,24}.

Most cases of COVID-19 have generally increased acute-phase reactants\textsuperscript{22}. Here, we focus on canonical and direct mediators of the systemic immune response secondary to an infection. All of the articles that reported C-reactive protein (CRP) showed increased levels\textsuperscript{2,3,7,9–12,14–16,19,21–24,27–29,31,34–37,41–43} except for one, whose cases were children and reported normal values\textsuperscript{40}. Additionally, serum ferritin and erythrocyte sedimentation rate (ESR) were above the normal reference values\textsuperscript{9–12,19,22,23,25,27,34,37}.

Unfortunately, procalcitonin levels reported among the reviewed articles did not allow us to establish a tendency since half of the articles reported normal levels\textsuperscript{1,2,5,22,36}, and the other half increased levels\textsuperscript{12,15,25,40} (Figure 2C).

Comparing SP with N-SP, increased levels of CRP in SP\textsuperscript{9,10,12,34} were reported in 80% of the references, while 100% reported increased levels of procalcitonin and ferritin in SP\textsuperscript{1,9,10,12,25,34,36}. In addition, 66% of the references reported normal levels of ESR independently of severity status\textsuperscript{10,11} (Figure 2D). Particularly, abnormally high procalcitonin levels in severe patients have been associated with the development of secondary infections\textsuperscript{1,27}. Although D-Dimer is not produced directly by the activation of the immune response, high levels of this marker in COVID-19 patients are strongly correlated with poor prognosis and, therefore, it has significant clinical value\textsuperscript{1,7,10,12,19,25,27,33}.

### Table 1. Characterization of the main titles.

| Immunological characteristics | Articles | Brief description | Type of articles |
|------------------------------|----------|------------------|-----------------|
| Clinical parameters associated with immune response | (1–3,5–7,9–43) | Description of cell number trends in CBC during COVID-19 and trends of acute phase reactants such as C reactive protein, procalcitonin, ferritin, etc. | Total:41 2 basic research, 4 commentaries, 12 observational, 9 case report, 10 reviews, 3 analytics, 1 clinical trial |
| Innate immune response | (1,3,6,7,9,10,12,14,15,20,23,24,31,33,41,44) | Description of the first line of defense against the SARS-Cov-2 | Total:16 4 Reviews, 3 commentaries, 5 descriptive, 2 case reports, 2 analytic |
| Cytokine storm | (1,7,9,10,12,23–26,34,36,45–48) | Identification of the cytokines implicated and the onset mechanism and development of this response | Total:15 4 Reviews, 1 commentary, 2 basic research, 7 descriptive, 1 analytic |
| Adaptive immunity | (1,3,6,7,9,10,12,18,20–22,24,31–33,41,42,44,45,47–53) | Descriptions and trends of T cells (CD4+ and CD8+), B cells and antibody changes | Total:26 3 basic research, 5 commentaries, 8 descriptive, 2 case report, 6 review, 2 analytic |
**Innate immune response**

Zhou et al. performed the identification and genomic characterization of SARS-CoV-2 and found that the virus uses angiotensin converting enzyme 2 (ACE2) and transmembrane serine protease 2 (TMPRSS2) as cell entry receptors. The ACE2 receptor is expressed on type II pneumocytes and immune cells such as monocytes, macrophages, and dendritic cells. In general, after a virus invades the host, it is initially recognized by the innate immune system through pattern recognition receptors (PRRs). The TLR3 and TLR7 recognize viral RNA, this event leads to recruitment of TRIF and MyD88 adapter molecules that activate NF-κB pathway, inducing the expression of type I interferons and the secretion of other proinflammatory cytokines for the activation of T cells, which limits the spreading of the virus. A well-coordinated innate and adaptive immune response may rapidly control the virus, while a failed immune response leads to viral spreading, cytokine storm, cytokine release syndrome (CRS), ARDS and death.

There is not enough knowledge of immunological indicators to understand the underlying mechanisms involved in COVID-19. Retrospective studies of SARS-CoV-2 infection have shown that there is limited information available on the host immune status, especially of the innate immune response. Besides the aforementioned parameters, SP compared to N-SP shared common features: 1. lower levels of natural killer (NK) cells and eosinophils in peripheral blood; 2. increased serum levels of IL-6 and TNFα; and 3. highly inflammatory macrophages with an enhanced chemokine production in the lung microenvironment.

In a cohort of 68 COVID-19 patients, the total number of NK cells decreased and had a reduced ability to produce CD107a, IFNγ, IL-2, GzmB and TNFα, and an upregulated expression of NKG2A, an inhibitor receptor characteristic of functional exhaustion. The decrease in NK cell count was even more pronounced in SP than in the N-SP group, but there was no significant difference in NK cell function. Moreover, SP had lower proportions of T and NK cells in bronchoalveolar lavage fluid (BALF) samples. Two groups have proposed different causes of low eosinophil count: Liu et al. proposed that lung damage due to SARS-CoV-2 may result in upregulation of glucocorticoid secretion, which in turn affects the production of eosinophils in the bone marrow; Du et al. argued that eosinophil depletion was related to lymphopenia since T cells produce IL-5, a cytokine that promotes eosinophil proliferation and activation in peripheral blood.

Monocyte accumulation resulted in elevated lung cytokine/chemokine levels, vascular leakage, and suboptimal T cell responses. A case report showed a lower frequency of CD16+CD14+ monocytes in peripheral blood of a positive case compared to a healthy donor (HD). Through gene expression analysis, Liao et al. identified monocyte-derived FCN1+ macrophages replacing the FABP4+ alveolar macrophages in the lung immune microenvironment of severe patients. These macrophages are potent chemokine producers as well as highly inflammatory.

**Cytokine storm**

Cytokine storm syndrome (CSS) is a form of systemic inflammatory response characterized by the excessive and uncontrolled release of a series of cytokines. CSS can be caused by a variety of disorders including infections and rheumatic diseases, and tumor immunotherapy. Clinically, it is commonly presented as systemic inflammation, ARDS, multiple organ failure and high inflammatory parameters.

Comparable to similar viral diseases such as SARS and MERS, most COVID-19 patients exhibit substantially elevated serum levels of both pro and anti-inflammatory cytokines. Some studies describe that IL-2, IL-6, IL-7, IL-10, G-CSF, IP-10, MCP-1, MIP-1A, and TNFα were higher and correlated with worse clinical progression and death. Levels of IL-6, IL-10, and TNFα correlated with T cell depletion. Other studies reported that additional cytokines and chemokines including IL-1β, sIL-1RA, CXCL8, IL-9, FGF, GM-CSF, IFNγ, G-CSF, PDGF, VEGF, IL-18, HGF, MCP-3, MIG, M-CSF, MIG-1a, CCL27 were increased as well.

In some studies, IL-6 was elevated in SP compared with N-SP throughout the clinical course and there was a correlation with disease severity. However, this was not consistent with the results obtained by other researchers, who did not find significant difference in serum levels of IL-6 between these groups. Interestingly, since IL-6 serum levels did not come from PBMCs, this cytokine might be produced by lung epithelial cells in COVID-19 patients.

Xiong et al. found that the expression of pro-inflammatory cytokines CXCL1, CXCL2, CXCL6, CXCL8, CXCL10/IP-10, CCL2/MCP-1, CCL3/MIP-1A, and CCL4/MIP1B was significantly upregulated in BALF samples of COVID-19 patients. Furthermore, increased transcription of respective chemokines receptors such as CCR2 (CCL2/MCP-1 receptor) and CCR5 (CCL3/MIP-1A receptor) was also observed, indicating the activation of these inflammatory signaling pathways. Additionally, elevated levels of macrophage chemoattractant CXCL10/IP-10, CCL2/MCP-1, neutrophil chemoattractant CXCL2, and CXCL8 facilitate the migration of these immune cells to the site of infection, which is consistent with mononuclear cell infiltrates observed in injured lung tissue.

Yang et al. compared the plasma cytokine/chemokine levels collected from SP and N-SP at different days after illness onset (d.a.o). The results showed that most of the cytokines were comparable between the two groups and maintained high levels, even at the later stage of the disease (≥ 15 d.a.o); however, IP-10, MCP-3, IL-1ra, and MIG were significantly higher in samples from SP. Further analysis showed that IP-10, MCP-3 and IL-1Ra expression levels were considerably high upon admission and were maintained during disease severity progression.
Figure 2. Clinical Parameters associated to Immune Response across references. 2A) Changes in CBC by citation percent: 79% reported leukocyte counts and 70% of them showed normal values, 23% reduced values, 3.3% increased values and 3.3% did not allow to establish a tendency. 100% reported lymphocyte counts and 63% of them showed reduced values, 29% normal values, 2.6% increased values and 5.3% did not allow to establish a tendency. 68.4% reported Neutrophil counts and 70% of them showed normal values, 13.8% increased values, 3.8% reduced values and 11.5% did not allow to establish a tendency. 13.1% reported Eosinophil counts, 80% of them showed decreased values and 20% normal values. 13.1% reported monocyte counts, 80% showed normal values and 20% decreased values. 10.5% reported basophil counts, 50% of them showed normal values and 50% increased values.

Figure 2B. Degree of change following disease severity in CBC and citation percent

Figure 2C. Levels of acute phase reactants by citation percent: 87.7% reported C-reactive protein and 93% of them reported increased levels, 3.5% normal levels and 3.5% did not allow to establish a tendency. 40.6% reported procalcitonin levels and 38.5% of them showed normal values, 38.5% increased levels and 23% did not allow to establish a tendency. 18.7% reported ferritin levels and 100% of them showed increased levels. 25% reported ESR levels and 75% of them showed increased levels, 12.5% normal levels and 12.5% did not allow to establish a tendency.

Figure 2D. Degree of change according to disease severity of acute phase reactants by citation percent. PCT: Procalcitonin CRP: C-reactive Protein, ESR: Erythrocyte sedimentation rate
and correlated with altered PaO2/FiO2 values and Murray scores. These results indicate that these three cytokines could be considered as predictive biomarkers of COVID-19 outcome\textsuperscript{26}. SARS-CoV-2 patients not only have upregulated inflammatory cytokines and chemokines that may lead to activated Th1 cell response, but also secreted excessive anti-inflammatory cytokines such as IL-10, IL-4, and TGF-\textbeta that may suppress inflammation by Th2 polarization\textsuperscript{26}. TGF-\textbeta signaling can be modulated by virus infection to prevent apoptosis and to promote fibroblast proliferation and myofibroblast differentiation, thus playing a critical role in the development of pulmonary fibrosis\textsuperscript{52}. These cytokines (pro and anti-inflammatory) mediate T cell depletion and extensive pulmonary pathology, leading to massive infiltration of neutrophils and macrophages, diffuse alveolar damage with the formation of hyaline membranes and diffuse thickening of the alveolar wall\textsuperscript{7}.

Elevated serum cytokine and chemokine levels in infected patients are correlated with disease severity and adverse outcome, suggesting a possible role for hyper-inflammatory responses in COVID-19 pathogenesis\textsuperscript{9,52}. All these data suggest that widespread lung damage associated with SARS-CoV-2 may be caused more by an exaggerated innate immune response than the virus itself. In table 2, we resume the principal findings.

**Adaptive immunity**

**T cell responses.** As mentioned above, several articles included in this review showed reports of lymphopenia in peripheral blood of COVID-19 patients\textsuperscript{10}, which correlated with disease severity\textsuperscript{9,10,12,18,51}. Additionally, this reduction seems to be age-dependent since elderly patients had lower counts than their young counterparts, which could be partially explaining the high morbidity and mortality rates within this population\textsuperscript{21,51}.

Surprisingly, autopsies of COVID-19 patients revealed destruction of secondary lymphoid organs and atrophy of the spleen\textsuperscript{9}. The immune infiltrate in the lungs of SP corresponded mainly to monocytes and macrophages, with a low presence of lymphocytes, unlike N-SP, whose BALF transcriptome showed high proportions of tissue-resident CD8 T cells (CD8 T cell)\textsuperscript{58}. These pathological findings suggest a “primary” cytokine storm, which is mainly produced from macrophages and epithelial cells, as opposed to a “secondary” storm that is produced by activated T cells\textsuperscript{59}, and are consistent with an upregulation of genes associated with an acute inflammatory response observed in epithelial cells from severe patients\textsuperscript{58}.

Within the cytokine storm caused by SARS-CoV-2 infection, high concentrations of IL-6, TNF\textalpha, and IL-10 have been reported, which could be causing the destruction of lymphatic nodes and depletion of tissue-resident and circulating lymphocytes\textsuperscript{9,12,24,49,51}. Both IL-6 and IL-10 can inhibit IFNy production and upregulate PD-1 expression in lymphocytes through SOCS-3 signaling\textsuperscript{6}, whilst TNF\textalpha can be recognized by TNFR1 and induce apoptosis by activating the p53 pathway, whose gene transcription profile was enriched in PBMCs from COVID-19 patients\textsuperscript{52}.

Supporting this theory, Diao et al. found that these three cytokines were negatively correlated with total lymphocyte counts and when serum levels of these cytokines subsided in the late stages of the disease, the number of lymphocytes increased. Moreover, high levels of IL-6 and increased expression of TNFR1 have been associated with aging, making elderly patients more susceptible to developing lymphopenia and therefore greater severity of the disease\textsuperscript{51}.

Another possible explanation for the lymphopenia observed in COVID-19 patients could be selective pyroptosis of these cells through the NLRP3 inflammasome activation induced by viral infection\textsuperscript{20,22}. Few studies concluded that the virus was not capable of infecting lymphocytes due to their low levels of ACE2 expression and because no viral RNA was detected in PBMCs\textsuperscript{52}. However, by using human T lymphocyte cell lines, researchers found that, unlike SARS-CoV, SARS-CoV-2 is capable of infecting these cells. This infection was receptor-dependent and mediated by the S protein of the virus, suggesting that ACE2 is not the only receptor that can interact with the virus\textsuperscript{52}. Other studies have proposed the possible action of CD147, CD163 and DC-SIGN, present in other leukocytes, as viral entry receptors\textsuperscript{52}. Fortunately, similar to MERS-CoV, SARS-CoV-2 infection appears to be abortive in lymphocytes, as the virus could not replicate after infection\textsuperscript{51}. Further studies are required to confirm whether this infection activates apoptotic pathways (i.e. pyroptosis) in lymphocytes, contributing to lower counts of immune cells.

Finally, reduction of circulating lymphocytes could also be associated with an impaired or limited lymphopoiesis caused by viral infection\textsuperscript{14} or redistribution of immune cells via...
attachment to the vascular endothelium or lung infiltration\textsuperscript{12}. Both of these hypotheses rely on the role of platelets in viral infection, as platelet-derived CXCL4 prevents Agglutinin-A from inhibiting lymphocyte generation in bone marrow, and thus when activated, they can enhance lymphocyte adhesion to the endothelium, thereby promoting homing to endothelial veins and migration to lungs\textsuperscript{31}.

To have an effective antiviral response, adequate activation of helper and cytotoxic lymphocytes is necessary. Hence, researchers have also characterized the frequencies, phenotypes, and functionality of lymphocytes subsets. CD4 T cell counts had the same behavior as total lymphocytes, with SP having the lowest numbers\textsuperscript{9,10,12,51}. Interestingly, a multiple linear regression model revealed that CD4 T cell counts could predict the duration of viral RNA in swabs and stool samples from patients\textsuperscript{42}.

The percentages of different CD4 T cell subsets also appear to influence the severity of the disease. A high proportion of naive CD4 T cell compartment is usually associated with a better response to infectious diseases. However, several authors reported a significant increase of this population in detriment of the CD45RO+ memory subset in SP\textsuperscript{31,7,9,48}. Isolated CD4 T cells from patients had a higher expression of TIGIT, an inhibitory receptor, and produced less IFNy, TNF\textalpha, and IL-2, cytokines characteristic of a Th1 response, when stimulated \textit{in vitro}; this phenomenon was even more noticeable in SP\textsuperscript{32}. In addition, the number of Tregs also seems to be lower in infected patients, especially in SP\textsuperscript{9,12}. In sum, a reduction of memory and regulatory subsets may contribute to the maintenance of a proinflammatory environment, which leads to exhaustion of T cells and an inadequate helper response.

CD8 T cell counts showed contradictory behaviors; Qin \textit{et al.} found no significant difference in the number of circulating CD8 between SP and N-SP. Nevertheless, they demonstrated that the number of CD8 T cells in COVID-19 patients was lower than in HD\textsuperscript{33}. In contrast, other researchers reported a significant decrease in CD8 T cells in SP compared to N-SP\textsuperscript{12,21,51}. In the lungs of N-SP, there was an increase of CD8 T cells, which expressed high levels of effector molecules including GzmA, GzmK, FAS-L, CCL5, and tissue residence markers such as ITGA1, CXCR6, and JAML\textsuperscript{48}. This cytotoxic CD8 infiltration may have an important role in virus elimination since these cells had an effector phenotype and a higher level of proliferation in N-SP that in SP. However, some authors found that the CD8 T cell population was also increased in SP lungs, which could be explained as a side effect of hyper-inflammation or late lung infiltrate\textsuperscript{48}.

In a case report of a COVID-19 patient, a peak of CD8+CD38+ HLA-DR+ cells, known for their role in antiviral response, was reported from day 7 to day 9, to finally fall on day 20\textsuperscript{41}. This subset was much higher than in HD, Zheng \textit{et al.} demonstrated that CD8 T cells from SP upregulate TIGIT and produce significantly lower amounts of GzmB and Perforin B than N-SP\textsuperscript{52}. Another study reported that CD8 T cells from COVID-19 patients expressed higher levels of PD-1 and that among them, SP had the highest percentages of CD8+PD1+ cells\textsuperscript{51}. This exhaustion phenotype in CD8 T cells was accompanied by a higher expression of NKG2A, with a lower expression of CD107a and lower production of IFNy, IL-2, and GzmB\textsuperscript{31}. In addition, SP had lower percentages of CD8+CD28+ cells, which could mean a decrease in the activation of these cells. However, Chen and Qin found no difference in the number of IFNy producing cells between patients and HD\textsuperscript{1}. Taken together, an improper T helper response accompanied by a strong inflammatory environment could result in the functional exhaustion of the cytotoxic compartment and therefore a progression of the disease, resulting in ARDS and subsequent death (Figure 4).

\textbf{B cell responses:} The production of neutralizing antibodies against SARS-CoV-2 plays an important role in the late stages of infection and might provide long-lasting memory that prevents reinfection\textsuperscript{31}. Qin and Chen found no difference in the total numbers of B cells between SP and N-SP\textsuperscript{9,12}. However, Chen reported differences in LB percentages between these two groups\textsuperscript{12}.

In a case report of one COVID-19 patient, the kinetics of antibody-secreting cells (ASCs) was described. ASCs appeared at the time of viral clearance (day 7), increasing significantly on day 9. The expansion of these cells was accompanied by an increase in circulating T follicular helper cells, which have an important role in isotype switch and affinity maturation of ASCs\textsuperscript{31}. Furthermore, when evaluating the transcriptional profile in PBMCs of patients with COVID-19, the signaling pathways related to humoral immune response mediated by circulating immunoglobulin and B cells immunity were upregulated\textsuperscript{12}.

\textbf{Antibodies:} Antibodies against SARS-CoV-2 were found in all patients samples one month after infection in both N-SP and SP subjects\textsuperscript{15,55,56}. Although no differences were found in either global antibodies or complement proteins between patients

| Cytokine | Non-Severe | Severe | References |
|----------|------------|--------|------------|
| TNFa | 3 | 11 | (1,9,12,51) |
| IL-1B | N/1 | N/1 | (1,12) |
| IL-1ra | 3 | 11 | (36) |
| IL-2R | N/1 | 3 | (12,36) |
| IL-6 | 3 | 11 | (10,12,25,34,36,51) |
| IL-8 | N/1 | N/1 | (1,12) |
| IL-10 | N/1 | N/1 | (1,36,51) |
| IFN\textgamma | 3 | 11 | (1,36,51) |
| IP10 | 3 | 11 | (1,36) |
| MCP1 | 3 | 11 | (1) |

N-SP: Non-Severe Patients SP: Severe Patients; N: normal; 1: Mildly Elevated; 11: Highly Elevated

\textbf{Table 2.} Comparison of serum cytokine concentrations between N-SP and SP in the studies included
and HD\textsuperscript{2}, Zhang et al. proposed that some patients had higher titers of antiphospholipid antibodies, including anticardiolipin and anti-B2-glycoprotein, which has been associated with the development of severe thrombosis and vascular injury\textsuperscript{24}.

The presence of anti-Nucleocapsid (NP) or anti-Receptor Binding Domain (RBD) IgG against SARS-CoV-2 in the serum of patients was correlated with neutralizing titers, therefore it is safe to conclude that neutralizing antibodies naturally develop in response to infection. The seroconversion of IgG and IgM of anti-RBD antibodies developed first than that of anti-NP and, surprisingly, the peak of seroconversion developed faster in SP. Another striking finding was that IgG seroconversion happened at the same time or even earlier than that of IgM, suggesting a dysregulated antibody production\textsuperscript{18,31,45}. IgM and IgG production reached a plateau on day 15 and day 21, respectively\textsuperscript{56}.

Higher production of antibodies is usually associated with a better response to infection, therefore, Kelvin et al assessed antibody titers in COVID-19 patients taking into account their age and comorbidities in order to investigate whether these could be affecting antibody production\textsuperscript{18}. Patients with or without comorbidities had similar levels of specific antibodies, and no significant difference was found between elderly and young adults, although the latter group always had a lower viral load during the disease. They also concluded that serum levels of these antibodies did not correlate with disease severity\textsuperscript{18}, however two other studies found an association between high levels of anti-SARS-CoV-2 antibodies and worse outcome of the disease\textsuperscript{7,55}. These last findings suggest the possibility of development of Antibody Dependent Enhancement during SARS-CoV-2 infection, indicating that SP could be producing non-neutralizing antibodies that overstimulate the innate immune system, increasing production of MCP1, IL-8, and IL-6 and thus aggravating the degree of inflammation and tissue damage, as well as exacerbating the previously mentioned lymphopenia\textsuperscript{7,53}.

**Question box: What immunological features of certain individuals allow them to be better equipped to resist or tolerate the infection?**

- Could prior infections reprogram the immune system, modifying the response to SARS-CoV-2?
- Could a tolerance-based approach rather than targeting the pathogen itself could lead to better treatments?
- What is the role of immunosenescence and inflamming in SARS-CoV-2 infection?
- Which are the best biomarkers associated with favorable prognosis?
- How does the immune profile changes, and to what extent in asymptomatic and recovered patients?
- How could the immune system adapt to the possible emergence of a mutated SARS-CoV-2 due to therapeutic pressure?
- Are there HLA molecules related to different outcomes?
- Are there any pulmonary morphologic characteristics that could contribute to an exacerbated immune response?
- Which mechanisms prevent SARS-CoV-2 replication within infected lymphocytes?
- Do the expression levels and functionality of the ACE2 receptor alter the immune response?

**Conclusions**

Immune response-associated clinical parameters such as lymphopenia, neutrophilia, and elevated levels of CRP and ferritin effectively correlates with disease severity and prognosis in SARS-CoV-2 infection. COVID-19 severe patients display immune dysregulation characterized by a hyperinflammatory response and sustained cytokine production that correlates with high mortality.
Moreover, disease severity in COVID-19 patients strongly correlated with lower T cells numbers accompanied by an overall lower Th1 cytokine production in CD4 cells. Concomitantly, increased expression of exhaustion markers and decreased secretion of effector cytokines in the CD8 compartment revealed an impaired cytotoxic response, especially in severe Patients. Considering that severe patients had a higher and faster rate of specific antibody production, a proper cellular compartment response seems to be more desirable for viral clearance.

Further research is needed to elucidate the immunological features that allow certain individuals to be better equipped to resist or tolerate the infection, therefore we propose some outstanding questions (Question BOX) to address this issue.

Limitations

Our scouring review has some limitations. The amount of original research continues increasing, and the quality of the research was not evaluated due to the recent development of the disease and the lack of original research. Reviews and comments rely on hypotheses, thus some of them must be proved in basic and clinical research.

Ethical disclosures

Funding. This research received no specific grant from any funding agency in the public, commercial, or not-for-profit sectors.

Conflicts of interest. The authors have no conflicts of interest.

References

1. Huang C, Wang Y, Li X, Ren L, Zhao J, Hu Y, et al. Clinical features of patients infected with 2019 novel coronavirus in Wuhan, China. Lancet. 15 de febrero de 2020;395(10223):497-506.
2. Liu K, Chen Y, Lin R, Han K. Clinical features of COVID-19 in elderly patients: A comparison with young and middle-aged patients. Journal of Infection [Internet]. 2020; Disponible en: http://www.embase.com/search/results?subaction=viewrecord&from=export&id=L2005431188 http://dx.doi.org/10.1016/j.jinf.2020.03.005
3. Cossarizza A, De Biasi S, Guaraldi G, Girardis M, Mussini C. SARS-CoV-2, the Virus that Causes COVID-19: Cytometry and the New Challenge for Analytical Cytology [Internet]. 2020; Disponible en: http://www.embase.com/search/results?subaction=viewrecord&from=export&id=L631282483 http://dx.doi.org/10.1002/cyto.a.24002
4. Report of the WHO-China Joint Mission on Coronavirus Disease 2019 (COVID-19) [Internet]. [citado 24 de marzo de 2020]. Disponible en: https://www.who.int/publications-detail/report-of-the-who-china-joint-mission-on-coronavirus-disease-2019-(covid-19)
5. Lai C-C, Liu YH, Wang C-Y, Wang Y-H, Hsueh S-C, Yen M-Y, et al. Dysregulation of immune response in patients with COVID-19 in Wuhan, China. Clinical infectious diseases : an official publication of the Infectious Diseases Society of America [Internet]. 2020; Disponible en: http://www.embase.com/search/results?subaction=viewrecord&from=export&id=L631358781 http://dx.doi.org/10.1122/jcic.103744
6. Cao X, Xiaoan C, Yanping C, Jian X, Xing Z, Sha X, et al. Risk Factors Associated with Acute Respiratory Distress Syndrome and Death in Patients with Coronavirus Disease 2019 Pneumonia in Wuhan, China. JAMA Internal Medicine [Internet]. 2020; Disponible en: http://www.embase.com/search/results?subaction=viewrecord&from=export&id=L631235378 http://dx.doi.org/10.1001/jamainternmed.2020.0994
7. Chen J, Qi T, Liu L, Ling Y, Qian Z, Li T, et al. Clinical progression of patients with COVID-19 in Shanghai, China. Journal of Infection. marzo de 2020;
8. Chen G, Wu D, Guo W, Cao Y, Huang D, Wang H, et al. Clinical and immunologic features in severe and moderate Coronavirus Disease 2019. The Journal of clinical investigation [Internet]. marzo de 2020; Disponible en: http://www.embase.com/search/results?subaction=viewrecord&from=export&id=L6311270519 http://dx.doi.org/10.1016/j.jcid.2020.03.013
9. Qin C, Zhou L, Hu Z, Zhang S, Yang S, Tao Y, et al. Disregulation of immune response in patients with COVID-19 in Wuhan, China. Clinical infectious diseases : an official publication of the Infectious Diseases Society of America [Internet]. 2020; Disponible en: http://www.embase.com/search/results?subaction=viewrecord&from=export&id=L631240468 http://dx.doi.org/10.1093/cid/ciaa248
10. Wu C, Xiaoan C, Yanping C, Jiana X, Xing Z, Sha X, et al. The deadly virus that causes COVID-19: Cytometry and the New Challenge for Analytical Cytology [Internet]. 2020; Disponible en: http://www.embase.com/search/results?subaction=viewrecord&from=export&id=L631379884 http://dx.doi.org/10.1016/j.cia.201711.0103
11. Haveri A, Smura T, Kuivenan S, Österlund P, Heipojoki J, Ikonen N, et al. Serological and molecular findings during SARS-CoV-2 infection: the first case study in Finland, January to February 2020. Euro surveillance : bulletin European sur les maladies transmissibles = European communicable disease bulletin. marzo de 2020;25(11)
12. Li X, Xu A, Zhang Y, Xuan W, Yan T, Pan K, et al. Patients of COVID-19 may benefit from sustained lopinavir–combined regimen and the increase of eosinophil may predict the outcome of COVID-19 progression. International journal of infectious diseases : IJID : official publication of the International Society for Infectious Diseases [Internet]. 2020; Disponible en: http://www.embase.com/search/results?subaction=viewrecord&from=export&id=L6311270519 http://dx.doi.org/10.1016/j.ijid.2020.03.013
13. Qu R, Ling Y, Zhang Y-H-Z, Li Y, Y-C, Li X-M, et al. Platelet-to-lymphocyte ratio is associated with prognosis in patients with coronavirus disease-19. Journal of Medical Virology [Internet]. 2020; Disponible en: http://www.embase.com/search/results?subaction=viewrecord&from=export&id=L2004505336 http://dx.doi.org/10.1002/jmv.25781
14. Jin X, Cao Y, Jiang N, Chen Y, Alwadi, X. Z, et al. Novel Coronavirus Pneumonia (COVID-19) Progression Course in 17 Discharged Patients: Comparison of Clinical and Thin-Section CT Features During Recovery. Clinical infectious diseases : an official publication of the Infectious Diseases Society of America [Internet]. 2020; Disponible en: http://www.embase.com/search/results?subaction=viewrecord&from=export&id=L631379884 http://dx.doi.org/10.1016/j.cia.201711.0103
15. Fu Y, Cheng Y, Wu Y. Understanding SARS-CoV-2-Mediated Inflammatory Responses: From Mechanisms to Potential Therapeutic Tools. Virolologia Sinica [Internet]. 2020; Disponible en: http://www.embase.com/search/results?subaction=viewrecord&from=export&id=L2004364468 http://dx.doi.org/10.1016/s12250-020-00207-4
16. Liu Y, Yang Y, Zhang C, Huang F, Wang F, Yuan J, et al. Clinical and biochemical indexes from 2019-nCoV infected patients linked to viral loads and lung injury. Science China Life sciences. 2020;63(3):364–374.
17. Yang Y, Peng F, Wang R, Guan K, Jiang T, Xu G, et al. The deadly coronaviruses: The 2003 SARS pandemic and the 2020 novel coronavirus pandemic. Journal of Autoimmunity. Academic Press; 2020.
18. Yi Y, Lagniton PNP, Ye S, Li E, Xu R-H. COVID-19: what has been learned and to be learned about the novel coronavirus disease. International Journal of Biological Sciences. marzo de 2020;16(10):1753–1766.
19. Zhang W, Zhao Y, Zhang F, Wang Q, Li T, Liu Z, et al. The use of anti-inflammatory drugs in the treatment of people with severe coronavirus
Human immune response to SARS-CoV-2: What is known? A scoping review

25. Zhou F, Yu T, Du R, Fan G, Liu Y, Li Z, et al. Clinical course and risk factors for mortality of adult inpatients with COVID-19 in Wuhan, China: A retrospective cohort study. The Lancet. marzo de 2020;395(10229):1054–1062.

26. Zhou G, Zhao Q. Perspectives on therapeutic neutralizing antibodies against the Novel Coronavirus SARS-CoV-2. Int J Biol Sci. 15 de marzo de 2020;16(10):1718-23.

27. Infantino M, Damiani A, Gobbi FL, Grossi V, Lari B, Macchia D, et al. Serological Assays for SARS-CoV-2: Infectious Disease: Benefits, Limitations and Perspectives. ISM Med Assoc. abril de 2020;22(4):203-10.

28. Leng Z, Zhu R, Hou W, Feng Y, Yang Y, Han Q, et al. Transplantation of ACE2(-) Mesenchymal Stem Cells Improves the Outcome of Patients with COVID-19 Pneumonia. Aging Dis. abril de 2020;11(2):216-28.

29. Rokni M, Ghasemi V, Tavakoli Z. Immune responses and pathogenesis of SARS-CoV-2 during an outbreak in Iran: Comparison with SARS and MERS. Reviews in medical virology. abril de 2020;

30. Chan JFW, Yuan S, Kok KH, To KKW, Chu H, Yang J, et al. A familial cluster of pneumonia associated with the 2019 novel coronavirus indicating person-to-person transmission: a study of a family cluster. The Lancet. febrero de 2020;395(10223):514-523.

31. Thevarajan I, Nguyen THO, Koutsakos M, Druce J, Caly L, van de Sandt CE, et al. Breadth of comitant immune responses prior to patient recovery: a case report of non-severe COVID-19. Nature Medicine [Internet]. 2020; Disponible en: http://www.embase.com/search/results?subaction=vie...f=export&id=L2004435182 http://dx.doi.org/10.1038/s41591-020-0819-2

32. Zheng H-Y, Zhang M, Yang C-X, Zhang N, Wang X-C, Yang X-P, et al. Elevated exhaustion levels and reduced functional diversity of T cells in peripheral blood may predict severe progression in COVID-19 patients. Cellular & Molecular Immunology [Internet]. 2020; Disponible en: http://www.embase.com/search/results?subaction=vie...f=export&id=L2004439947 http://dx.doi.org/10.1038/s41423-020-0401-3

33. Zheng M, Gao Y, Wang G, Song G, Liu S, Sun D, et al. Functional exhaustion of antiviral lymphocytes in COVID-19 patients. Cellular & Molecular Immunology [Internet]. marzo de 2020; Disponible en: http://www.embase.com/search/results?subaction=vie...f=export&id=L631330101 http://dx.doi.org/10.1038/s41423-020-0402-2

34. Ruan Q, Yang K, Wang W, Jiang L, Song J. Correction to: Clinical predictors of mortality due to COVID-19 based on an analysis of data of 150 patients from Wuhan, China. Intensive Care Medicine [Internet]. abril de 2020; Disponible en: http://link.springer.com/10.1007/s00134-020-06028-2

35. Zhang X, Cai H, Hu J, Lian J, Gu J, Zhang S, et al. Epidemiological, clinical characteristics of cases of SARS-CoV-2 infection with abnormal imaging findings. International journal of infectious diseases : IJID : official publication of the International Society for Infectious Diseases [Internet]. abril de 2020; Disponible en: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4174766/

36. Yang Y, Shen C, Li J, Yuan J, Yang M, Wang F, et al. Exuberant elevation of IP-10, MCP-3 and IL-1ra during SARS-CoV-2 infection is associated with disease severity and fatal outcome. medRxiv. marzo de 2020;2019;2020.03.02.20029975.

37. Cheng Z, Lu Y, Cao Q, Qin L, Pan Z, Yan F, et al. Clinical Features and Chest CT Manifestations of Coronavirus Disease 2019 (COVID-19) in a Single-Center Study in Shanghai, China. AJR American journal of roentgenology. 2020;1-6.

38. Huang W-H, Teng L-C, Yeh T-K, Chen Y-J, Lo W-J, Wu M-J, et al. 2019 novel coronavirus disease (COVID-19) in Taiwan: Reports of two cases from Wuhan. J Microbiol Immunol Infect. 19 de febrero de 2020;

39. Lee N-Y, Li CW, Tsai H-P, Chen P-L, Syue L-S, Li M-C, et al. A case of COVID-19 and pneumonia returning from Macau in Taiwan: Clinical course and anti-SARS-CoV-2 IgG dynamic. J Microbiol Immunol Infect. 10 de marzo de 2020;

40. Xing Y-H, Ni W, Wu Q, Li W-J, Li G-J, Wang W-D, et al. Prolonged viral shedding in feces of pediatric patients with coronavirus disease 2019. J Microbiol Immunol Infect. 28 de marzo de 2020;

41. Promptetchara E, Ketloy C, Palaga T. Immune responses in COVID-19 and potential vaccines: Lessons learned from SARS and MERS epidemic. Asian Pacific journal of allergy and immunology. 2020;38(1):1–9.

42. Ling Y, Xu SB, Lin YX, Tian D, Zhu ZQ, Dai FH, et al. Persistence and clearance of viral RNA in 2019 novel coronavirus disease rehabilitation patients. Chinese medical journal. febrero de 2020;

43. Xu Y-H, Dong J-H, An W-M, Lu X-Y, Yin X-P, Zhang J-Z, et al. Clinical and computed tomographic imaging features of novel coronavirus pneumonia caused by SARS-CoV-2. J Infect. abril de 2020;80(4):394-400.

44. Hirano T, Murakami M. COVID-19: A New Virus, but a Familiar Receptor and Cytokine Release Syndrome. Immunity. 19 de abril de 2020;

45. Zhou P, Yang X-L, Wang X-G, Hu B, Zhang L, Zhang W, et al. A pneumonia outbreak associated with a new coronavirus of probable bat origin. Nature. mayo de 2020;579(7798):270–273.

46. Moore JB. June CH. Cytokine release syndrome in severe COVID-19. Science. 1 de mayo de 2020;368(6490):473-4.

47. Kawasaki T, Kawai T. Toll-Like Receptor Signaling Pathways. Front Immunol [Internet]. 25 de septiembre de 2014 [citado 8 de mayo de 2020];5. Disponible en: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4174766/

48. Liao M, Yuan L, Yuan J, Wen Y, Xu G, Zhao J, et al. The landscape of lung bronchoalveolar immune cells in COVID-19 revealed by single-cell RNA sequencing. medRxiv. febrero de 2020;2020.02.23.20026690.

49. Sarzi-Puttini P, Giorgi V, Sinotti S, Marotto D, Ardizzone S, Rizzardi G, et al. COVID-19, cytokines and immunosuppression: what can we learn from severe acute respiratory syndrome? Clinical and experimental rheumatology [Internet]. marzo de 2020; Disponible en: http://www.ncbi.nlm.nih.gov/pubmed/32202240

50. Hyun-Jung Lee C, Koohy H. In sihouette identification of Vaccine targets for 2019-nCoV. F1000Res. 2020.9;145.

51. Diao B, Wang C, Tan Y, Chen X, Liu Y, Ning L, et al. Reduction and Functional Exhaustion of T Cells in Patients with Coronavirus Disease 2019 (COVID-19). medRxiv. 20 de febrero de 2020;2020.02.18.20024364.

52. Xiong Y, Yuan L, Liu C, Dehe W, Guo M, Jiang A, et al. Transcriptomic characteristics of bronchoalveolar lavage fluid and peripheral blood mononuclear cells in COVID-19 patients. Emerging Microbes and Infections. 2020;9(1):761–770.

53. Wang X, Xu W, Hu G, Xia S, Sun Z, Liu Z, et al. SARS-CoV-2 infects T lymphocytes through its spike protein-mediated membrane fusion. Cell Mol Immunol. 7 de abril de 2020;

54. Vellingiri B, Jayaramayya K, Iyer M, Narayanamas Y, Govindasamy A, Giridharan B, et al. COVID-19: A promising cure for the global panic. Sci Total Environ. 4 de abril de 2020;725:138277.

55. Zhao J, Yuan Q, Wang H, Liu W, Liao X, Su Y, et al. Antibody responses to SARS-CoV-2 in patients of novel coronavirus disease 2019. Clinical Infectious Diseases [Internet]. marzo de 2020; Disponible en: http://www.embase.com/search/results?subaction=vie...f=export&id=L631367949 http://dx.doi.org/10.1093/cid/ciaa344 https://academic.oup.com/cid/advance-article/doi/10.1093/cid/ciaa344/5812996

56. Guo L, Ren L, Yang S, Xiao M, Chang D, Yang F, et al. Profiling Early Humoral Response to Diagnose Novel Coronavirus Disease (COVID-19). Clinical infectious diseases : an official publication of the Infectious Diseases Society of America [Internet]. marzo de 2020; Disponible en: http://www.embase.com/search/results?subaction=vie...f=export&id=L631299351 http://dx.doi.org/10.1093/cid/ciaa310