Gut microbiome, Vitamin D, ACE2 interactions are critical factors in immune-senescence and inflammaging: key for vaccine response and severity of COVID-19 infection

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Abstract
Background The SARS-CoV-2 pandemic continues to spread sporadically in the Unites States and worldwide. The severity and mortality excessively affected the frail elderly with co-existing medical diseases. There is growing evidence that cross-talk between the gut microbiome, Vitamin D and RAS/ACE2 system is essential for a balanced functioning of the elderly immune system and in regulating inflammation. In this review, we hypothesize that the state of gut microbiome, prior to infection determines the outcome associated with COVID-19 sepsis and may also be a critical factor in success to vaccination.

Methods Articles from PubMed/Medline searches were reviewed using a combination of terms “SARS-CoV-2, COVID-19, Inflammaging, Immune-senescence, Gut microbiome, Vitamin D, RAS/ACE2, Vaccination”.

Conclusion Evidence indicates a complex association between gut microbiota, ACE-2 expression and Vitamin D in COVID-19 severity. Status of gut microbiome is highly predictive of the blood molecular signatures and inflammatory markers and host responses to infection. Vitamin D has immunomodulatory function in innate and adaptive immune responses to viral infection. Anti-inflammatory functions of Vit D include regulation of gut microbiome and maintaining microbial diversity. It promotes growth of gut-friendly commensal strains of Bifida and Fermicuttus species. In addition, Vitamin D is a negative regulator for expression of renin and interacts with the RAS/ACE/ACE-2 signaling axis. Collectively, this triad may be the critical, link in determination of outcomes in SARS-CoV-2 infection. The presented data are empirical and informative. Further research using advanced systems biology techniques and artificial intelligence-assisted integration could assist with correlation of the gut microbiome with sepsis and vaccine responses. Modulating these factors may impact in guiding the success of vaccines and clinical outcomes in COVID-19 infections.

Keywords COVID-19 sepsis · Gut microbiome · Vitamin D · ACE2 · Immune-senescence · Inflammaging · Vaccination

Introduction
The SARS-CoV-2 pandemic continues to spread sporadically in the Unites States and worldwide. We are one year into the pandemic and there is no clear therapy, and the global vaccination of the population is in its early stages and too soon to determine any long-term efficacy in the vaccinated population. To complicate matters further, there is appearance of breakthrough infections with mutant strains of SARS-CoV-2 globally, with high infectiousness and varying severity [1, 2]. This current COVID-19 pandemic has established that our understanding of the pathophysiology of sepsis remains incomplete and further research is needed to fill these gaps. It has been demonstrated that combinations of risk factors, such as medical comorbidities, older age group, mitochondrial dysfunction, along with high nasopharyngeal viral load, are predictors for worse outcomes [3, 4].

There is a wide heterogeneity in COVID-19 disease severity, ranging from asymptomatic to fatal disease. However, it is well established that elderly patients are susceptible to severe sepsis and death compared with younger population. Even among the elderly, it is the frail patients with co-existing illnesses, signifying advanced biological age rather than chronological age which influences this susceptibility [5].
Immune-senescence and inflamming are natural phenomenon associated with aging. It defines frailty and biological susceptibility to COVID-19 sepsis and its outcomes. Immune senescence is described as age-related remodeling of immune cell repertoire and functions in the human physiology. Inflamming represents a state of chronic inflammation, elevated cytokine levels, associated with inflammasome activation and aging [6]. Both immune-senescence and inflamming are complex interrelated systemic processes and encompass both circulating cells and tissue immune system, such as mucosal lining of respiratory and gastrointestinal (GI) tract, resulting in changes, for example in the microbiome. The GI tract has a long surface area and in addition to its absorptive, metabolic functions also harbors a robust immune system and function [7]. There is growing evidence that cross-talk between gut microbiome and immune system is vital for priming, development, and function of immune cells, both locally and systemically. This can affect the overall health of an organism [8]. Gut immune system is associated with severity of lung infections, pneumonia and sepsis through gut lung axis [9, 10]. Dysbiosis is defined as an alteration in the composition of gut microbiome or the production of abnormal microbial peptides that may disrupt the natural symbiotic harmony between the host and microbiome with deleterious consequences. Dysbiosis along with immune-senescence and inflamming leads to progressive reduction in the ability to trigger effective antibody and cellular responses against infections and vaccinations. This is demonstrated in seasonal respiratory tract viruses, such as influenza infections and vaccinations [6, 11, 12].

There appears to be a complex interaction between gut microbiota, ACE-2 expression and Vitamin D in COVID-19 pathogenesis. This triad may provide the critical, missing link in determination of outcomes in SARS-CoV-2 infection (Fig. 1). We hypothesize that the state of gut microbiome, prior to infection determines the outcome associated with COVID-19 sepsis. In this review, we present several lines of evidence and discuss how inflamming, immune-senescence and immune health of gut may affect the severity of COVID-19 sepsis and responses to vaccination. Simple, but strategic nutritional, non-pharmacologic interventions may immune-modulate the elderly immune system and mitigate risks and severity of COVID-19 infections. An in-depth understanding of these mechanisms will have an impact in guiding the success of vaccines and discovery of novel diagnostic and therapeutic targets which are needed against novel SARS-CoV-2 virus.

**Impact on innate immunity**

There are consequences of immune-senescence in COVID-19 sepsis. Changes in composition of functional DC cells, dysfunctional NK cell cytotoxicity and loss of naïve T cells are critical in COVID-19 infections. There is a delay in type I interferon (IFN) activation and together with dysfunctional NK cell-mediated cytotoxicity impedes viral clearance. Early type I IFNs is crucial for restricting viral replication and dissemination, through autocrine and paracrine type I IFN receptor (IFNAR) signaling [11–13, 18, 19].

DC cells are antigen-presenting cells (APCs). They express toll-like receptors (TLRs) that recognize conserved pathogen-associated molecular patterns (PAMPs) on microbes and are key regulators in antimicrobial host defense. Recognition of antiviral components by TLRs is vital to the secretion of type I IFN and cytokines that facilitate the coordination of innate antiviral immunity [11, 12, 18]. Subsequently DCs present these peptides through MHC II (major histocompatibility) program to T and B lymphocytes, thus stimulating adaptive and humoral immune responses, respectively. The functions of DCs are impaired in elderly with delayed maturation and migration demonstrated during microbial invasion [13]. Experiments have demonstrated that DCs from lymph nodes in adult mice have diminished ability to prime T cells compared to DCs from young mice. This impaired function is due to...
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translocation of JAK/STAT1 pathways [23]. Degradation of intracellular IFN and delay of early antiviral defense IFN response, juxtaposed with elevated cytokine, and subsequent exaggerated T cell response define the hyper-inflammation features of COVID-19 [13, 19, 24, 25].

This hyper-inflammatory response was demonstrated in critically ill patients with COVID-19 where-in an impaired early interferon (IFN) type I response is characterized by lack of IFN-β, IFN-α production and activity. These patients exhibited persistent viremia and an exacerbated inflammatory response [25, 26]. Lei et al. demonstrated that SARS-CoV-2 viral proteins: NSP1, NSP3, NSP12, NSP13, NSP14, ORF3, ORF6 and M protein inhibit IFN-β promoter activation and act as intracytoplasmic IFN antagonist. This delay in intracellular antiviral response enables the virus to replicate rapidly. Large amounts of viral transcripts with high viral loads were observed before the IFN induction in SARS-CoV-2-infected cells [22]. Therefore, lack of timely and robust early antiviral response is central in COVID-19 pathogenesis [22, 25, 26].

**Impact on the adaptive immunity**

Similar to its effects on innate immunity, immune-senescence and inflamming also impact the adaptive immune system. The thymus begins to involute with onset of puberty and there is significant shift in T cell immune profile with aging. There is a relative paucity of naïve T cells with abundance in differentiated memory T and B cells. This predisposes elderly to a blunted antibody and antiviral response, especially to novel pathogens and antigens. The decrease in naïve cells seems related to changes in sex hormones and to decrease of IL-7, a hematopoietic growth factor secreted by stromal cells in bone marrow and thymus [12].

Decrease in T cell diversity with fewer naïve T cells, reversal of CD4+/CD8+ cell ratio and increase in differentiated memory T cell population signify immune senescence. Further chronic metabolic diseases, chronic viral infections shift the immune system toward an inflammatory, autoimmune, profile [12, 13, 20].

CD4+ T cell profile demonstrates significant changes with aging. The CD4+ profile consists of Th1, Th2, Th17 and Treg (Regulatory T cells). Th1/CD4+ cells are pro-inflammatory and secrete IFN-γ, IL-1, and IL-2. Th2/CD4+ cells are anti-inflammatory and secrete IL-4, IL-10 that tempers down the Th1 response to prevent uncontrolled damage due to cytokine storm. In a normal physiological state, the Th1/Th2 ratio is low and Th2 anti-inflammatory cells are upregulated. This is observed in healthy elderly patients [13, 17]. However, in the presence of an infection, a smooth transition to upregulate Th1 profile is necessary, to eliminate invading viruses and other pathogens but minimize tissue destruction [17]. Dysregulated function in this switch in the elderly leads to the classical cytokine storm observed in the COVID-19 sepsis [13, 27]. SARS-CoV-2 infection has demonstrated lymphopenia associated with severe sepsis. In most COVID-19 patients, there is a decrease in CD4+ and CD8+ T cell counts and in CD4+/CD8+ ratio. In addition, T cells have demonstrated signs of exhaustion [27].

Experiments on mice have demonstrated the interaction of respiratory viral infections, such as influenza virus with gut microbial and immune system through gut lung axis. Lung-derived CD4+ T cell destroys microbiota homeostasis and promotes resident Th17 cell polarization. Gut epithelium produces chemokines CCL25/CCR9 that mediates the recruitment of lung-derived CD4+ T cell into the small intestine. CCL25–CCR9 axis contributes to altering the composition of the intestinal microbiota after influenza infection and development of intestinal inflammation by recruiting these effector lymphocytes into the intestinal mucosa [28]. Elderly are particularly susceptible to these effects as there is upregulated pro-inflammatory Th17 cells with loss of anti-inflammatory TREG activity associated with immune senescence [13].

Age-related immune senescence also affects B cell repertoire. There is reduced diversity and number of B cells and receptors as well as immunoglobulin isotypes and a decrease in specific humoral immune responses against new extra-cellular pathogens. There is a shift from immunoglobulin produced by naïve cells (IgD, IgM) to immunoglobulin produced by memory B cells (IgG, IgA) [12]. Impaired functions of DC together with loss of naïve T cells are primary factors for suboptimal immune response to viral infections in elderly [12, 29].

Other important contributor to immune senescence includes loss of telomerase activity with telomere shortening in aged T cells [12, 17, 30]. T cells upregulate telomerase activity to compensate for telomere loss, incurred during proliferation in response infections and chronic inflammatory conditions. However, repeated antigen exposure as seen with chronic antigenic stimulation by Herpes and CMV viruses leads to clonal expansion of memory T cell phenotype. This results in telomere shortening leading to replicative senescence. These senescent, exhausted T cells do not express the co-stimulatory molecule CD28, required for activation of T cells and also express immune check point inhibitors such as PD-1 and the cytotoxic T-lymphocyte antigen (CTLA)-4 [12, 13, 15, 17, 30–32]. T cell exhaustion and anergy and its positive correlation with severity of sepsis are demonstrated in SARS-CoV-2 infections. T cells from COVID-19 patients have significantly higher levels of exhausted marker, such as PD-1 and Tim-3 expression [27, 33].

Chronic medical conditions associated with aging, such as diabetes, hyperlipidemia, and chronic inflammatory diseases lead to inflamming. Chronic chemical, physical,
and nutritional antigentic stimulation also trigger oxidative stress and inflammation. Inflamming represents a state of chronically elevated cytokine levels and associated inflammasome activation. The markers for inflamming include TNF-α, C-reactive protein, IL-6, NF-κB and are chronically elevated in the elderly [34]. Mitochondrial dysfunction with increased reactive oxygen species (ROS), inefficient oxidative-phosphorylation and loss of mitochondrial antiviral signaling (MAVS) are associated with aging and may contribute to immune-senescence and inflamming [4, 17, 35, 36]. Impaired autophagy of damaged cells, elevated reactive oxygen species (ROS) and elevated levels of cellular dysfunctional proteins result in further activation of TLRs and inflammasomes and thus a chronic inflammatory state [13, 17, 20].

Collectively, this evidence suggests that aging-associated immune-senescence and inflamming contributes to a diminished and altered response by the innate and adaptive immunities to acute viral infections such as novel SARS-CoV-2 infections [12, 13, 15, 18, 30–32].

**Gut immune system**

The GI tract has an elaborate, dynamic, complex immune system called gut-associated lymphoid tissue (GALT). It is interspersed with lymphatic tissues called Peyer’s patches and extensive sub-epithelial immune cell repertoire and mesenteric lymph nodes [37]. This complex immune environment is constantly exposed to challenges arising, both in the lumen and systemic internal environment [7]. There is normal physiological mucosal turn over and shedding, intermittent luminal trauma from ingested food, loss of integrity of cell junctions which releases additional antigens. The immune cells constantly sample food antigens, pathobiont, such as other non-commensal bacteria, viruses and other foreign peptides [37, 38]. The gut epithelium along with the immune cells produces protective factors, such as secretory IgA, α-Defensins, antimicrobial peptides and mucus. Immune cells in gut remain vigilant through complex interactions and partake in a delicate balance between homeostasis, tolerance and inflammation. Tipping this balance may lead to increased risk for sepsis (translocation, growth of pathobiont) or excessive immune destruction (autoimmunity) [7, 8, 39].

Systemic immune-senescence and inflamming also have damaging effects on the GI tract and dysfunctional gut immune system is associated with severity of lung infections, pneumonia and sepsis through gut lung axis [9, 15, 28]. The state of gut function and its physiological health, such as dysbiosis, vitamin D deficiency, poor nutrition, obesity, along with COVID-19-related factors, such as viral load, may determine the outcome associated with COVID-19 sepsis [3, 12, 18].

GI tract also expresses ACE-2 receptors, and it is confirmed that SARS-CoV-2 virus commonly infects the GI tract epithelium. The duration of viral replication varies greatly, and prolonged shedding of viral RNA in stool samples has been reported after resolution of symptoms [40–42]. In the subsequent sections, we discuss the interaction between dysbiosis, ACE-2 expression and Vitamin D with the gut immune system, and its possible association with severity in COVID-19 sepsis.

**Complex interaction between microbiota, ACE-2 expression and Vitamin D in COVID pathogenesis: the missing link: (Fig. 1)**

Healthy microbiome is essential in development and balancing the function of both pro- and anti-inflammatory T-cell pathways [8, 43, 44]. Baseline dysbiosis is demonstrated to have a major impact on the local and systemic immunity by altering the gastrointestinal tract environment [45]. Aging is associated with frailty, malnutrition, generalized atherosclerosis resulting in decreased mucosal blood flow and villous atrophy. Intestinal dysbiosis shifts the intestinal microbiota from obligate anaerobes, such as Bifidobacterium spp., Lactobacillus spp., and Faecalibacterium prausnitzii to facultative anaerobes, such as pathogenic Streptococci, Staphylococci, Enterococci and Enterobacteria [10, 15]. Reduced biodiversity associated with chronic antibiotic use, inflammatory diseases and aging may result in colonization of toxin-producing Clostridium difficile bacteria. These changes lead to an alteration of gut immune cell composition creating a state of low-grade chronic inflammation. In addition, there is loss of protective factors, such as mucus, immunoglobulins and other antimicrobial protective peptides, such as bacteriocins. Dysbiosis is also implicated with other chronic diseases associated with aging. These include increased susceptibility to respiratory infections, inflammatory bowel disease, chronic inflammation, arthritis and also malignancy of the bowel [15, 46–48].

In addition, intestinal dysbiosis is also implicated in progression of malignancy and resistance to anticancer immunotherapy. These effects occur due to modulation of the immune system [49, 50]. Further dysbiosis, through diet and aging and associated medical diseases, such as COPD, chronic inflammation and antibiotics, are linked with altered immune responses and homeostasis in the airways [51].

Induction of a pro-inflammatory T-cell response by microorganisms can elicit development of T-helper-1 (Th1), Th2, or Th17 cells [28]. Pro-inflammatory T-cell helper and cytotoxic responses are suppressed by the action of various subsets of regulatory T cells (Treg) [13, 52]. Loss of
gut mucosal membrane integrity, loss of tight junction and leaky gut phenomenon lead to activation of local PAMP molecules. Pattern recognition receptors, such as TLR, NOD recognize these molecules and activate inflammasomes with IL-6, IL-17, TNF release, causing oxidative stress and release of ROS [52, 53].

There is evidence that dysbiosis may be relevant in systemic SARS-CoV-2 infections.

Yeoh et al. demonstrated in their cohort of 100 patients that composition of gut microbiome in patients with COVID-19 correlated with disease severity and plasma concentrations of several inflammatory cytokines, chemokines associated with tissue damage (TNF-α, CXCL10, CCL2 and IL-10, C-reactive protein, lactate dehydrogenase, aspartate aminotransferase and gamma-glutamyl transferase.). Patients with COVID-19 were depleted of beneficial microbiome with immunomodulatory potential, such as Faecalibacterium prausnitzii, Eubacterium rectale and several Bifidobacterium species and the dysbiosis persisted after the clearance of the virus [54].

In another cohort, Gu et al. compared healthy controls with COVID-19 patients by 16S rDNA gene sequencing of fecal samples. COVID-19 patients had a significantly reduced bacterial diversity with higher abundance of opportunistic pathogens, such as Streptococcus, Rothia, Veillonella and Actinomyces, and a lower relative abundance of beneficial symbionts. These patients also had a significantly higher level of IL-6 and TNF-α, CRP and significantly lower lymphocyte count, markers for severity and mortality [55].

Similarly, Khan et al. demonstrated an association of dysbiosis with severe inflammatory response. Suppressed Firmicutes/Bacteroidetes ratio, caused by the depletion of the fiber-utilizing bacteria F. prausnitzii, B. Plebius, and Prevotella, and a relative increase in Bacteroidetes species is associated elevated serum IL-21 levels. However, they did not elaborate on the temporal association of the inflammatory markers with the stool samples. Elevated IL-21 level early in viral infections is beneficial and is cytotoxic to the infected cells, however, a delayed response is detrimental as observed in COVID-19 infections [56].

In an NIH-funded study, Venzon et al. demonstrate a link between SARS-CoV-2 infection, gut microbiome dysbiosis, and bloodstream infections. The loss of microbiome diversity and Faecalibacterium in patients with severe COVID-19 infection mirrored a similar loss of diversity in mice inoculated with high doses of SARS-CoV-2. Stool samples of 101 COVID-19 patients at two different clinical sites revealed substantial gut microbiome dysbiosis, paralleling observations in the animal model. This analysis revealed that genus Faecalibacterium was negatively associated with blood stream infection (OR = 1.49, CI [−2.82, −0.18]). Analysis of blood culture results with paired microbiome data obtained from these patients suggests that bacteria translocate from the gut into the systemic circulation of COVID-19 patients [57].

Tang et al. in their cohort of 57 patients with COVID-19 with varying severity indicated that dysbiosis occurred in COVID-19 patients and changes in the gut microbiome was associated with disease severity and hematological parameters. The butyrate-producing bacteria, Faecalibacterium prausnitzii, Clostridium butyricum, Clostridium leptum, and Eubacterium rectale, decreased significantly in severe patients. This was accompanied by an increase in the opportunistic pathogens Enterococcus, Enterobacteriaceae [58].

Similarly, Moreira-Rosário et al. in their cohort on 115 patients demonstrated the gut microbiota of moderate and severe patients have a lower Firmicutes/Bacteroidetes ratio, predominance of Proteobacteria and lower abundance of beneficial butyrate-producing bacteria. Multivariable regression analysis showed that the Shannon diversity index [odds ratio (OR) 2.85, 95% CI 1.09–7.41, p = 0.032] and C-reactive protein (OR 3.45, 95% CI 1.33–8.91, p = 0.011) are risk factors for severe COVID-19 [59].

Collectively, these data suggest that loss of butyrate-producing obligate anaerobes in the gut may increase the severity of COVID-19-related sepsis and involve a state of systemic hyper-inflammation with its un-intended consequences, such as generalized hypoxia and mucosal ischemia. Sepsis causes structural and functional changes in the mucosa of the GI tract, such as loss of mucus, barrier integrity, change in microbiome and risk of translocation. In addition, there may be direct mucosal SARS-CoV-2 viral infection through the blood viremia or swallowed virus. As mentioned earlier, SARS-CoV-2 has been detected in feces implicating either an active infection or asymptomatic viral shedding [40, 41, 60]. This systemic hyper-inflammatory state when combined with dysfunction of GI tract may overwhelm the body’s ability to recover from cytokine storm.

Therefore, a healthy baseline microbiota of the GI tract may be necessary and may have beneficial effects in a population if they get infected with SARS-CoV-2 virus. The mechanism by which gut microbiota modulates the immune system is varied and includes a combination of pro-inflammatory and anti-inflammatory function. Gut microbiome produce diverse metabolites from the anaerobic fermentation of exogenous undigested dietary components. One such group of molecules is short-chain fatty acids (SCFAs)—such as butyrate, propionic acid and acetic acid. These are produced by microbial fermentation of undigested or partially digested dietary fiber and have broad effects on epithelial barrier and host immune system function [8, 61]. It promotes production of mucus, secretion of secretory IgA to enhance local immunity. SCFA also inhibits NF-kB activity and downstream activation of inflammasomes and cytokines [61]. There is upregulation of colonic regulatory T (Treg) cells and their production of anti-inflammatory cytokines,
such as transforming growth factor-β (TGFβ) and IL-10, to maintain a delicate balance between homeostasis and tolerance [52, 53, 61–63]. Other functions of SCFA include activating and facilitating migration of DC cells at sites of inflammation and infection. Activated DC cells as described earlier, are vital as an antiviral defense mechanism by releasing IFN in COVID-19 sepsis [52, 53, 61, 62]. SCFA activities also effect cell metabolism and may also inhibit transcription of genes by inhibiting histone deacetylase (HDAC) activity [63]. SCFA activity is also implicated in gut–lung axis as these microbial products may circulate systemically to the airway epithelium and alter inflammation locally in airway epithelium. Increased circulating levels of SCFAs lead to enhanced generation of dendritic cell precursors through bone marrow hematopoiesis and subsequent seeding of lungs by DCs with high antiviral and primary defense capacity [64].

Other molecules derived from microbiome with anti-inflammatory activity in the GI tract are sphingolipids, cerebromides and polysaccharides. Microbiome-derived sphingolipids, demonstrate stimulation of DC to increase expansion of virus-specific T-cells upon viral infections [65]. Polysaccharide A (PSA) suppresses inflammation by upregulating activity of colonic regulatory T (Treg) cells, and boosting their production of anti-inflammatory cytokines [52, 66].

Other effects of gut dysbiosis include loss of mucosal integrity and antecedent risk for translocation of lipopolysaccharides (LPS) which are derived from pathogenic Gram-negative bacteria. This commonly occurs in sepsis with generalized tissue hypo-perfusion leading to gut mucosal ischemia and loss of barrier function. LPS triggers activation of toll-like receptor (TLR4) and its downstream signaling pathways manifested as a cytokine storm [67, 68].

There is sufficient evidence that probiotics defined as live non-pathogenic, microbial ingredients when consumed in sufficient quantities, have health benefits. They have immunomodulatory properties and are strain-dependent. For example, introduction of probiotic strains, such as *Bifidobacterium lactis*, results in a significant increase in the proportion of mononuclear leukocytes, and enhances the activity of NK cells [10, 15, 69] Majority of probiotics consist of *Lactobacillus, Bifidobacterium, Saccharomyces* and may offer protection in COVID-19 sepsis.

Zhang et al. reported a propensity score-matched retrospective study evaluating probiotics and outcomes in 375 adult patients with COVID-19, 179 cases (probiotics group) and the other 196 cases (non-probiotics group) were included. The primary outcome was clinical improvement which was compared among propensity-score-matched groups and an unmatched cohort. Secondary outcomes included the duration of viral shedding, fever, and hospital stay. Among the propensity-score-matched groups, probiotics was related to clinical improvement (log-rank \( p = 0.028 \)). This relationship was driven primarily by a shorter (days) time to clinical improvement [difference, \(-3\) \((-4\) to \(-1\)], \( p = 0.022 \), reduction in duration of fever [\(-1.0\) \((-2.0\) to \(0.0\)], \( p = 0.025\), viral shedding [\(-3\) \((-6\) to \(-1\)], \( p < 0.001\), and hospital stay [\(-3\) \((-5\) to \(-1\)], \( p = 0.009\). However, the limitations were, this study was a retrospective propensity-matched and not a prospective study, single institution in Shenzhen and critically ill patients were underrepresented [70].

A recent pilot study on fifteen patients by Zhou et al., shed insight into association between intestinal dysbiosis and GI epithelial ACE-2 levels and severity of COVID-19 [71]. Increased *Coprobacillus* and pathogenic *Clostridium* species with concomitant decrease in *Faecalibacterium prausnitzii* correlated with increased severity of COVID-19 sepsis. Another observation of this study was that *Bacteroides* species downregulated the expression of angiotensin-converting enzyme 2 (ACE2) and correlated inversely with SARS-CoV-2 load in fecal samples from patients [71].

It has been demonstrated that SARS-CoV-2 binds to human angiotensin-converting enzyme 2 (ACE-2) to infect the host cell and is vital step for entry into the cell [72, 73]. In addition to virus entry, this process also leads to downregulation of anti-inflammatory ACE2 expression, thereby causing excessive generation of pro-inflammatory Angiotensin II via the enzyme ACE [74]. The renin–angiotensin system (RAS) is a critical homeostasis regulatory system in the human physiology. The ACE-Angiotensin II-AT1R pathway is called the classical RAS axis and is vital for various normal physiological functions, while the ACE2-Angiotensin1-7-MasR signaling is called the counter-regulatory RAS axis pathway, and is equally important for negative regulatory role and anti-inflammatory activity [75, 76]. The primary role of RAS axis is to stimulate the sympathetic nervous system causing vasoconstriction and maintain the vascular tone and blood pressure. However, with chronic overstimulation, they cause unintended deleterious side effects of inflammation, oxidative stress, atherosclerosis, migration of endothelial cells and vascular smooth muscle cells, fibrosis, and myocardial hypertrophy. The negative regulatory axis mediated by ACE2 can antagonize these effects. ACE2 levels decline with aging and so does its protective effect on many of organ systems [70, 76, 77].

The interaction between dysbiosis and ACE-2 enzyme activity in GI tract remains poorly understood. ACE-2 is expressed on the epithelium of GI tract and also other epithelial surfaces [78]. Deficiencies in murine (ACE-2) knock out studies have demonstrated increased susceptibility to dysbiosis and intestinal inflammation and colitis, induced by epithelial damage [79]. Fecal transplantation of the altered microbiota from mutant ACE-2 knock out mice into germ-free wild-type was able to transmit this increased susceptibility to develop severe colitis. Hashimoto et al.
demonstrated the possible anti-inflammatory role of ACE-2 enzyme activity in GI tract through amino acid hemostasis. ACE2-dependent changes in epithelial immunity and gut microbiota can be directly regulated by dietary amino acid tryptophan [80]. Similarly, other authors have also suggested an RAS-independent protective mechanism of ACE-2 expression, such as alteration in intestinal amino acid transporter function, tryptophan uptake, expression of antimicrobial peptides, and changing the ecology of the gut microbiome [79, 81]. Elderly patients with decreased ACE-2 levels and deregulated (overactive) RAS axis as depicted with hypertension, diabetes and kidney disease have increased risk for dysbiosis and this incidentally remains as a factor in elevated morbidity and mortality associated with COVID-19 sepsis [70, 76, 77].

Vit D, in addition to its function in bone and mineral metabolism has an essential immunomodulatory function in innate and adaptive immune responses to viral infection [82]. Anti-inflammatory functions of Vit D include regulation of gut microbiome and maintaining microbial diversity. Vit D promotes growth of gut-friendly commensal strains of Bifida and Fermentisus species. Vit D and its receptor (VDR) deficiency results in dysbiosis, leaky guts resulting in chronic, low-grade inflammation in the gastrointestinal tract and is implicated as an etiology in inflammatory bowel disease [83, 84]. In addition, Vit D also regulates microbial complexity, maintains cell junctions and barrier function and the mucosal immune responses to ensure intestinal homeostasis [83]. Vit D along with other nutrient factors acts in synergy to regulate ZO-1, occludin and claudin tight junction proteins. These are key proteins essential for the integrity and intact intestinal barrier [83].

Vit D plays a key role in innate antiviral and a subsequent anti-inflammatory (immune tolerance) phase in the COVID-19 sepsis [77, 83, 85, 86]. Vit D is essential in attenuating cytokine storm by decreasing the TNF-α/IFN-β and IFN-γ, IL-6 activity and also modulating adaptive immunity by suppressing T helper cell type 1 response and promoting T regulatory cell function [83, 84]. In Vit D-deficient states, there is an increased ratio of cytotoxic T lymphocytes and reduced TREG lymphocytes.

In addition, Vit D is a negative regulator for expression of renin and its production and interacts with the RAS/ACE/ACE-2 signaling axis. Vit D deficiency leads to an upregulation of RAS/ACE signaling and pro-inflammatory activity. This may enhance the severity in COVID-19 sepsis with clinical features of ARDS, pulmonary edema, cardiovascular injury and thrombotic events [77, 87].

Immune cells have a key role in the synthesis of active form of Vit D and are also a target for its effects. They activate the hormone and aid the local immune function in an autocrine or a paracrine fashion. There is evidence that antigen-presenting cells (APC), such as macrophages and dendritic cells, synthesize the active form of Vit D, 1,25-dihydroxyvitamin D (1,25 (OH)2D) from its precursor 25-hydroxyvitamin D (25-OHD) via the enzyme 1α-hydroxylase (CYP27B1) [85, 86]. Active form of Vit D and its receptor expression are regulated during differentiation of human monocytes into macrophages and may be crucial for effective function of the primary antigen-presenting immune cells against invading pathogens [85, 88].

Normal level of active forms of Vit D levels is critically dependent upon 1α-hydroxylase (CYP27B1), a mitochondrial cytochrome P450 enzyme that catalyzes the conversion of inactive precursor of Vit D to active metabolite 1α25(OH)2D3. This 1-hydroxylase enzymatic activity is localized to mitochondrial inner membranes and may decline in immune-senescent cells [89, 90]. In addition to the immune cells, the enzyme 1α-hydroxylase (CYP27B1) is also expressed in the epithelium of the respiratory and GI tract [90].

Vit D, thus has an essential role in complex interaction in maintaining GI homeostasis, gut microbiome, RAS/ACE/ACE-2 signaling, differentiation and function of immune cells and innate, humoral, and cellular immune responses. The beneficial effects of Vit D supplementation in viral infections remain unclear. A prior pooled meta-analysis study of 11,321 participants from 25 randomized controlled trials demonstrated that Vit D supplementation protected against acute respiratory tract infections. Oral supplementation in patients with very low serum levels (< 25 nmol/L) of 25-hydroxyvitamin D concentrations, was demonstrated as safe and protected against acute respiratory tract infection [91]. The recent COVID-19 pandemic has demonstrated the high morbidity and mortality in the elderly patients and the minorities (dark skinned) in the countries of northern latitude compared to similar population in the southern latitudes. This observational and circumstantial evidence has suggested that long winters, remaining indoors, less exposure to sunlight and inadequate absorption of Vit D from the sunlight leads to deficiencies and may remain an essential factor [92].

Although promising the evidence of Vitamin D supplementation in acute COVID-19 infection remains inconsistent and insufficient in reducing the probability of ICU admission, inflammation, hospitalization, and pulmonary involvement. In a metaanalysis by Kazemi et al. on 39 retrospective and prospective cohort, cross-sectional, case-control, and randomized controlled trial studies to assess the relation between 25D status and SARS-CoV-2 infection as well as COVID-19 severity. Overall, the researchers noted a greater risk of SARS-CoV-2 infection in the Vitamin D-deficient group. However, the studies were heterogeneous in methodological and statistical approach, and with many confounding variables, such as age, sex and ethnicity, diet, sunshine,
lifestyle choices and caution should be exercised in interpreting these results [93].

In addition, a randomized controlled trial on 240 COVID-19 patients with moderate to severe COVID-19, a single oral dose of 200,000 IU of vitamin D₃, compared to placebo, did not significantly reduce the duration of hospitalization (median of 7.0 vs. 7.0 days; unadjusted hazard ratio for hospital discharge, 1.07) and did not support the use of a high dose of vitamin D₃ for treatment of moderate to severe infection [94]. Currently, a large number of trials on benefits of Vitamin D in COVID-19 are being conducted at multiple international centers. Their details can be accessed at www.clinicaltrials.gov and results are awaited.

There also is some evidence that Vit D may have salutary influence in response to immunization. However, most studies reporting on the favorable effects of Vit D on vaccine responses have investigated responses to influenza vaccination and the results are conflicting [95]. Patients with renal failure on hemodialysis receiving parenteral calcitriol treatment may respond with higher antibody responses for influenza, Hepatitis B and tetanus vaccines [95].

**Inflamming, immune-senescence, immune health of the gut and vaccination in elderly**

Efficacy and robustness of vaccinations are suboptimal in elderly compared to children and younger population. As described, immuno-senescence and inflamming may collectively diminish the immune responses and ability to fight infections in elderly. There is growing evidence that age-associated immune-senescence and inflamming also have a detrimental effect on vaccine response in elderly [6, 18, 96]. Aging and decline in naïve T cells may be risk factors for failure to generate a coordinated immune response, resulting in increased susceptibility to severe COVID-19 [97]. Elderly patients also have a rapid decline in antibodies to vaccinations and frequently need a booster dose to sustain levels of antibodies. For example, the ability of influenza vaccine to induce protection is age-related, with an efficacy between 70 and 90% in children and young adults, but reduces to 30–50% for those over 65 years [11, 98]. Profiles predicting successful vaccination response to influenza vaccination, correlated with positive expression of genes associated with T-cell and B-cell function while monocyte (TLR4, TLR8, NOD2, and ASGR2) pattern recognition receptors involved in innate immunity and inflammation-related genes encoding (IFN-γ, IL-13R, TIMP2, LYN, SYK) negatively correlated with influenza-specific antibody responses, supporting the concept that inflammatory responses (inflamming and immune-senescence) at baseline might be detrimental to vaccine-induced antibody responses [14, 45, 99].

In addition, older patients with immune-senescence lymphocytes may fail to mount a robust antibody response to newer virus and vaccines [19, 96] A clue to this possibility has emerged with recent reports of COVID-19 recurrences in a few patients getting re-infected by the variants of virus and due to waning of natural antibody response [1, 2, 100, 101]. Some of the re-infected patients suffered mild symptoms but other presented with severe symptoms on reinfec tion. This discrepancy and likely variation in immunological memory is a cause of concern in the vaccine and antibody therapies. Age-related changes in antigen uptake by APCs, loss of naïve lymphocytes, processing and presentation to MHC, as well as recognition and functional defects of B and T cells, may lead to reduced antibody responses [19]. This problem is especially worrisome in the setting of current SARS-CoV-2 pandemic, wherein the majorities of patients are elderly and have demonstrated the highest mortality regardless of comorbidities or ethnicity. Although promising with greater than 90% efficacy, the currently available COVID-19 vaccine-induced antibody response may be short-lived and wane in the elderly and remains to be noted.

The immunomodulatory property of probiotics may influence the response to vaccines [15, 43, 44]. A meta-analysis of nine RCTs including 623 participants, suggested that probiotics and prebiotics are effective in elevating immunogenicity by influencing seroconversion and protection rates in adults administered influenza vaccines. Participants who took probiotics or prebiotics showed significant improvements in the H1N1 strain protection rate [odds ratio (OR) 1.83, 95% confidence interval (CI) 1.19–2.82, \( p = 0.006 \)], the H3N2 strain protection rate (OR 2.85, 95% CI 1.59–5.10, \( p < 0.001 \)) and the B strain seroconversion rate (OR 2.11, 95% CI 1.38–3.21, \( p < 0.001 \)) [102]. Another meta-analysis evaluated 3812 patients from 26 different studies found a beneficial effect of probiotics. The evidence for a beneficial effect of probiotics on vaccine response was strongest for oral vaccinations for non-respiratory infections and for parenteral influenza vaccination [103]. The gut-associated lymphoid tissues (GALTs) and complex interactions with microbiota may thus play an important role in the induction of antigen-specific immune responses in the gut [37, 104].

**Future directions**

This review suggest that, until there are effective antiviral medications for COVID-19 infections, simple adjunct interventions that modulate immune systems, such as use of probiotics and Vit D, may offer protection to prevent severe sepsis in elderly and susceptible individuals. The underlying mechanisms can be harnessed for developing therapies that may offer some immune protection. The gut mucosal immune system along with its interaction through gut–lung axis may also be used to develop mucosal vaccines and can also be a candidate site for future vaccinations [44, 45, 104].
These adjunct therapies may improve vaccine efficacy and duration of protection for vaccinated individuals. However, there are certain challenges which need to be addressed. With advanced next-generation sequence technologies, such as meta-taxonomic, metabolomics, metagenomics and transcriptome analysis, along with multi-parametric high throughput flow cytometry, the gut microbiome can be characterized, and their resulting host humoral and cellular immune responses can also be studied. Advanced system biology and AI (artificial intelligence)-assisted integration could assist with correlation of the gut microbiome with vaccine responses [44]. Using machine learning models, Guo et al. demonstrated in their multi-omics analyses, a risk score based on 20 blood proteomic biomarkers, predicting severity of COVID-19 infection. They demonstrated that a core set of gut microbe can predict the proteomic biomarkers among 301 individuals and these correlated with pro-inflammatory cytokines in another independent set of 366 individuals [105]. Fecal metabolomics analysis suggests potential amino acid-related pathways linking gut microbiota to host metabolism and inflammation. Similar data may be needed in other ethnic, sex and age groups. Further studies are needed on establishing optimal probiotic strains, doses of probiotics and Vit D and timing of administration in relation to disease and vaccinations.

**Conclusion**

To conclude, SARS-CoV-2 pandemic continues to spread erratically and has caused destruction in terms of the economy, health and wellbeing of the population. We are one year into the pandemic and there seems to be no clear therapy. Although there is hope that vaccinations may control this pandemic eventually, the majority of global population is currently un-immunized, and it is too early to determine any long-term protective antibody levels and efficacy. We have described the important functions of gut microbiome, Vit D, and ACE-2/RAS system interaction in COVID-19 infection. Modifying these factors may attenuate the severity of illness in infected population. Further research is needed in role of probiotics in maintaining long-term immunity and success with vaccines in SARS-CoV-2 infection.

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