

## HYPOTHESES

# Sialylation of host proteins as targetable risk factor for COVID-19 susceptibility and spreading: A hypothesis

Antonella Bongiovanni<sup>1</sup> | Antonella Cusimano<sup>1</sup> | Ida Annunziata<sup>2</sup> | Alessandra d'Azzo<sup>2</sup> 

<sup>1</sup>Institute of Biomedical Research and Innovation (IRIB), National Research Council (CNR), Palermo, Italy

<sup>2</sup>Department of Genetics, St. Jude Children's Research Hospital, Memphis, Tennessee, USA

### Correspondence

Alessandra d'Azzo, Department of Genetics, St. Jude Children's Research Hospital, 262 Danny Thomas Place, Memphis, TN 38105, USA.  
Email: sandra.dazzo@stjude.org

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### Abstract

Individuals infected with the severe acute respiratory syndrome (SARS)-related coronavirus 2 (SARS-CoV-2) develop a critical and even fatal disease, called Coronavirus disease-19 (COVID-19), that eventually evolves into acute respiratory distress syndrome. The gravity of the SARS-CoV-2 pandemic, the escalating number of confirmed cases around the world, the many unknowns related to the virus mode of action, and the heterogenous outcome of COVID-19 disease in the population ask for the rapid development of alternative approaches, including repurposing of existing drugs, that may dampen virus infectivity. SARS-CoV-2 infects human cells through interaction with sialylated receptors at the surface of epithelial cells, such as angiotensin-converting enzyme 2 (ACE2). Glycan composition on virus entry receptors has been shown to influence the rate of infection of SARS-CoV-2 and spreading of virions has recently been linked to altered lysosomal exocytosis. These processes could concurrently involve the lysosomal system and its glycosidases. We hypothesize that modulating the activity of one of them, the lysosomal sialidase NEU1, could impinge on both the sialylation status of ACE2 and other host receptors as well as the extent of lysosomal exocytosis. Thus NEU1-controlled pathways may represent therapeutic targets, which could impact on SARS-CoV-2 susceptibility, infectivity, and spread.

### KEYWORDS

ACE2, COVID-19, lysosomal sialidase, NEU1, SARS-CoV-2, sialylation, spike protein

## 1 | INTRODUCTION

Sialylation is a crucial posttranslational modification in mammalian cells, because it affects, among others, the half-life, structural conformation, and ligand recognition of glycoconjugates.<sup>1</sup> This is particularly important in the field of host/pathogen interactions, where sialic acids are exploited by hosts to evade recognition by pathogens and by pathogens to escape the immune system response.<sup>2,3</sup> Given their terminal positions on glycan chains and electronegative charge, sialic acids influence the structural and biochemical properties

of a wide range of sialo-glycoconjugates, and, consequently, serve as important determinants in basic processes, including innate immunity, inflammation and viral recognition.<sup>4</sup> It is, therefore, not surprising that changes in the sialic acid content of host cell receptors as well as viral glycoproteins influence the tropism, pathogenicity, and infectivity of several viruses, including the novel SARS-CoV-2.

As of November 2020, SARS-CoV-2 has caused 58.425.681 confirmed cases of COVID-19, including 1.385.218 deaths (<https://covid19.who.int>) worldwide. The clinical features of patients, with confirmed diagnosis

of COVID-19, include lower respiratory tract illness with fever, dry cough, and dyspnea, manifestations similar to those seen in two other diseases caused by coronaviruses, SARS and Middle East respiratory syndrome (MERS).<sup>5</sup> It is now established, that these symptoms are accompanied by a fulminant and damaging immune reaction, referred to as the “cytokine storm,” and involve not only the lungs but also multiple other organs, including the heart and the brain.<sup>6-8</sup> With regard to the severe lung disease that ensues upon SARS-CoV-2 infection, the most common lung manifestations in COVID-19 patients are bilateral ground-glass opacity with subpleural distribution and absence of pleural effusion, a typical lung injury caused by viral pneumonia, as previously seen in SARS and MERS.<sup>9</sup> Following the 2003 epidemic of SARS, it was noticed that many patients, who survived the severe illness, developed residual pulmonary fibrosis, as shown by clinical findings and radiography.<sup>10</sup> Varying degrees of fibrosis were also observed in autopsies of COVID-19 fatal cases, and they appear to be a common feature in patients infected by SARS-coronaviruses.<sup>10</sup>

Based on data derived from the general population, it is clear that the pathogenesis and clinical outcome of coronavirus infection tend to be influenced by the age of the patients. The disease seems to affect more males than females, and older individuals with preexisting conditions, like hypertension, diabetes, obesity, and other cardiovascular diseases, have an increased risk of developing severe COVID-19.<sup>11</sup> Despite this age-related trend, SARS-CoV-2 infection does not spare young individuals, suggesting that genetic and/or epigenetic predisposition mechanisms might also play an important role. Here, we hypothesize that one such mechanism may involve the activity of the lysosomal sialidase NEU1. By modulating the levels of sialic acids on ACE2 (and other entry receptors), NEU1 may directly influence virus entry. We further postulate that NEU1 may contribute to virus spreading by directly controlling the process of lysosomal exocytosis.

## 1.1 | SARS-CoV-2 entry receptors: role of sialylation

Similar to other coronaviruses, SARS-CoV-2 has been shown to infect host cells through interaction between the receptor binding domain (RBD) of the trimeric S1 subunit of its spike (S) glycoprotein (S1 and S2 subunit composition) and ACE2 on the surface of human epithelial cells of the lung, heart, kidney, intestine, and brain.<sup>12,13</sup> ACE2 is a sialylated, single-pass, type I transmembrane enzyme that belongs to the angiotensin-converting enzyme family of dipeptidyl carboxydipeptidases.<sup>14</sup> The external enzymatic domain of ACE2 binds SARS-CoV-2 prior to the fusion of

the viral envelope with the host PM and endocytosis of the virus. This virus-receptor interaction requires priming of the spike S1 protein by the host serine protease TMPRSS2 and furin.<sup>12,15</sup> Structural studies using cryo-electron microscopy have shown that human ACE2, in complex with the amino acid transporter B<sup>0</sup>AT1, exists as a dimer and as such interacts with the RBD of the viral S glycoprotein of SARS-CoV-2.<sup>13</sup> Structural alignment of this receptor-virus complex suggests that two S protein trimers can simultaneously bind to an ACE2 homodimer.<sup>13</sup> It is noteworthy that, besides its functional domains, also the glycans on ACE2, and in particular its sialic acids, play an active role in the infection process.<sup>16,17</sup>

In addition to ACE2, Sigrist et al. have suggested that SARS-CoV-2 utilizes integrins (including  $\beta$ 1- and  $\beta$ 3-integrins) as cell surface receptors in one or more host species; in this case, attachment of the virus to these receptors occurs through a conserved Arg-Gly-Asp motif embedded in the RBD of all SARS-CoV-2 that have been sequenced until now.<sup>18</sup> Given that the functions of integrins are known to be influenced by their glycosylation and sialylation status,<sup>19</sup> it is likely that these posttranslational modifications also impact virus binding and, in turn, infection efficacy. Recently, another glycosylated and sialylated membrane protein, Neuropilin-1 (NRP1) has been shown to function as SARS-CoV-2 entry receptor, enhancing virus infectivity.<sup>20,21</sup> Furin cleavage of SARS-CoV-2 S1 protein allows binding to NRP1; subsequent processing of furin-cleaved S1 by cathepsins and TMPRSS2 allows virus entry and infection.<sup>20,21</sup> These results suggest that the interaction between the viral spike protein and host NRP1 may represent a potential target for antiviral therapy.

Lastly, studies of MERS have proposed that the MERS-CoV spike glycoprotein, which recognizes and binds to the dipeptidyl peptidase 4 (DPP4) entry receptor on respiratory epithelial cells, also uses sialic acids exposed at the cell surface as viral entry sites. In the case of MERS-CoV the sialic acid binding site has been identified and found that it is homologous to a region upstream of the RBD of the SARS-CoV-2 spike protein (the highly variable region at the N-terminus, NTD).<sup>22-24</sup> Thus, not only the functional activity of the entry receptors on the host cells but also the extent and type of their glycan chains, in particular their sialic acids, exposed at the cell surface may influence and modulate the rate of infection of the SARS-CoV-2, as demonstrated for other members of this virus family.<sup>17,25,26</sup>

## 1.2 | Chloroquine and hydroxychloroquine, pros and cons

Among the drugs used to treat SARS-CoV-2 infection, one well-known antimalaria drug, chloroquine, and its less toxic hydroxychloroquine, have gained a lot of attention

at the beginning of the pandemic but raised concerns and controversies later on.<sup>27-37</sup> When added in cultured cells, the non-protonated portion of chloroquine enters the cell, where it becomes protonated and concentrated in acidic, low-pH organelles, such as Golgi vesicles, endosomes, and lysosomes.<sup>38</sup> Intra-vesicular pH is known to control basic cellular functions, including *N*-glycosylation trimming/modification, organellar trafficking, and enzymatic activities. By increasing the pH of Golgi vesicles, chloroquine (and also NH<sub>4</sub>Cl and possibly hydroxychloroquine) is thought to interfere with terminal glycosylation, specifically sialylation, of the Golgi form of ACE2, likely by inhibiting Golgi sialyltransferases,<sup>39</sup> without affecting ACE2 expression levels at the cell surface.<sup>40-43</sup> In line with this finding is a recent modeling study showing that the sialylated portion of cell surface gangliosides, which functions as co-receptor for ACE2 and binds to the NTD of the spike glycoprotein of SARS-CoV-2, is blocked by both chloroquine and hydroxychloroquine.<sup>44</sup> This hinders the host-cell entry and subsequent virus replication, and, in turn, inhibits SARS-CoV-2 infection.<sup>41-43</sup> However, an opposite scenario could also be envisaged based on the effects of chloroquine on raising the lysosomal pH,<sup>45</sup> which inhibits the activity of lysosomal hydrolases, including NEU1. The latter could prevent desialylation of SARS-CoV-2 receptors that would recycle to the PM in their sialylated state and facilitate viral entry. These two opposing effects which may depend on the drug regimen and time of administration could explain the contradictory response to chloroquine/hydroxychloroquine treatment observed in COVID-19 patients.

Collectively these examples point to sialylation as an important factor to be considered for the development of potential new treatments for COVID-19 and for the design of new vaccines.

### 1.3 | Multifactorial roles of the lysosomal sialidase NEU1

In eukaryotes, processing/removal of sialic acids on sialo-glycoconjugates occurs via the action of 4 sialidases or neuraminidases, NEU1-NEU4, that are differentially expressed in cells and tissues, and have a different pH optimum and substrate specificity.<sup>46</sup> These enzymes have been shown to participate in and regulate many biological and pathological processes, from cell-cell communication and cell/tissue development and differentiation to cancer progression and metastasis.<sup>47,48</sup> Of the 4 mammalian sialidases, NEU1 is the most abundant and ubiquitously expressed,<sup>49</sup> and it is the only sialidase known to date that is deficient in two lysosomal storage diseases, sialidosis (NEU1 deficiency) and galactosialidosis (cathepsin A primary defect and secondary

deficiency of NEU1).<sup>49,50</sup> These pediatric conditions, associated with severe systemic and neurological abnormalities, share clinical manifestations with common adult diseases of aging, such as Alzheimer's disease, diabetes, atherosclerosis, obesity, fibrosis, and cancer.<sup>51-53</sup>

NEU1 catalyzes the removal of  $\alpha$ 2,3- or  $\alpha$ 2,6-linked sialic acids from the non-reducing end of glycan chains on sialo-glycoconjugates, with a preference for sialo-glycoproteins, an activity that is exerted at an acidic pH optimum. NEU1 localizes primarily in the endosomes/lysosomes but it is also found in the PM; both these cellular compartments are sites of desialylation of glycoproteins by NEU1.<sup>49,54,55</sup> Deficient or defective enzyme activity results in impaired processing/degradation of sialo-glycoproteins, and in their accumulation in a sialylated state, a feature that alters their biochemical properties and function, and is at the basis of pathogenesis in sialidosis.<sup>48,49,51,53,56-58</sup> However, the pathological consequences of impaired desialylation downstream of NEU1 loss of function depend on the type of substrates the enzyme targets *in vivo* and on their subcellular distribution, which, in turn, affect specific cellular pathways.<sup>48,49,51,57,59-66</sup> For instance, by processing the sialic acids on the lysosomal associated membrane protein 1 (LAMP1), NEU1 negatively regulates the calcium-dependent, physiological process of lysosomal exocytosis.<sup>57</sup> In cells with deficient or downregulated NEU1, lysosomes decorated with a sialylated, long-lived, LAMP1 tend to preferentially dock at the PM, ready to engage in lysosomal exocytosis upon changes in calcium concentration.<sup>57</sup> The end result is excessive release of lysosomal contents, including soluble hydrolases and exosomes, into the extracellular space with deleterious consequences for PM and extracellular matrix integrity.<sup>48,51,53,56-58</sup> Importantly, in fibroblasts from sialidosis patients with different degrees of disease severity, the residual NEU1 activity inversely correlated with the levels of LAMP1 redistributed at the PM, identifying this feature as indicative of the extent of lysosomal exocytosis.<sup>57</sup>

Based on this collective evidence, we hypothesize that reduced NEU1 activity in cells could have a dual effect: it could affect SARS-COV-2 infectivity because of impaired processing of the sialic acids in entry receptors and could promote virus release via LAMP1-dependent, excessive lysosomal exocytosis. In support of this hypothesis is the recent finding that  $\beta$ -coronaviruses use lysosomal exocytosis to egress into the extracellular environment.<sup>67</sup> These authors showed that the viral genome was enriched in lysosomes and that, upon viral infection, LAMP1 levels increased at the PM. They also demonstrated a de-acidification of lysosomes, which affected the activity of lysosomal enzymes and inhibited antigen presentation by the host cells.<sup>67</sup>

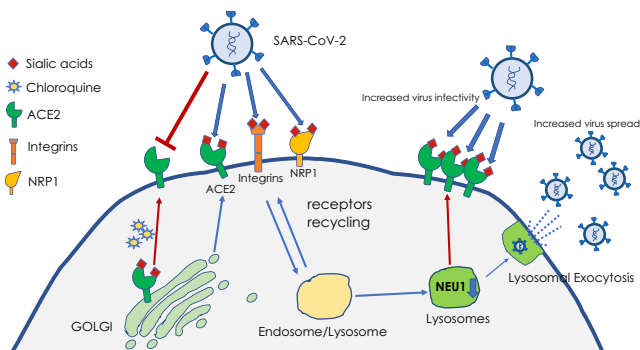
Lastly, a major downstream effect of NEU1 deficiency seen in the mouse model of sialidosis is a relentless expansion of the connective tissue, ultimately resulting in generalized

fibrosis.<sup>53</sup> The underlying pathogenic process leading to this fibrotic disease is initiated and perpetuated by the excessive exocytosis by NEU1-deficient myofibroblasts of soluble lysosomal contents and exosomes loaded with pro-fibrotic signaling molecules, like TGF $\beta$ .<sup>53</sup> The fact that NEU1 is downregulated in a cohort of patient-derived idiopathic pulmonary fibrosis (IPF) fibroblasts suggests a role for NEU1 deficiency/downregulation in causing or hastening the course of a fibrotic disease in the adult human population. It is, therefore, tempting to speculate that systemic downregulation of NEU1 coupled to excessive lysosomal exocytosis could represent a risk factor for the initiation and/or progression of the fibrotic disease associated with SARS-CoV-2. This is also supported by the failure of neuraminidase inhibitors, such as oseltamivir, peramivir, and zanamivir in the treatment of COVID-19 patients,<sup>68,69</sup> while those are first-line medications for the treatment and prophylaxis of influenza.<sup>70,71</sup>

## 2 | HYPOTHESIS

The premises outlined above have led us to design a workflow (Figure 1) aimed to identify additional functional biomarkers that will help to further characterize susceptibility to SARS-CoV-2 infection in the general population. We hypothesize that NEU1 levels inversely correlate with the extent of sialylation of the ACE2 receptor (and other entry receptors) and, in turn, the predisposition for entry of SARS-CoV-2 (Figure 1).

Furthermore, low NEU1 levels may also result in LAMP1-mediated excessive lysosomal exocytosis of lysosomal contents/exosomes and SARS-CoV-2 virions. Thus, NEU1 levels may also predict the tendency of an individual to become a SARS-CoV-2 “super spreader” (Figure 1).



**FIGURE 1** Proposed model of viral entry and spread. Low NEU1 activity results in excessive lysosomal exocytosis of viral particles and impaired desialylation of host receptors (ie, ACE2) recycled to the PM. These combined effects may lead to the higher infectivity and spreading of the SARS-CoV-2 virions. In contrast, chloroquine can block ACE2 sialylation in the Golgi, resulting in the exposure of undersialylated receptors at the PM and decreased susceptibility to the virus

In this context, it would be conceivable to:

1. IDENTIFY A RISK FACTOR FOR SARS-CoV-2 SUSCEPTIBILITY and SPREAD
  - by evaluating the expression levels of *NEU1* mRNA in human samples (the lowest NEU1 levels should correspond to worst prognosis and fit our super spreader hypothesis).
  - by screening the general population for *NEU1* synonymous variants known to lower NEU1 enzyme activity<sup>72</sup> as well as pathogenic mutations in carriers. These analyses will ascertain the involvement of NEU1 functional variants in the pathogenesis of COVID-19.
2. APPLY UNCONVENTIONAL THERAPY TO INCREASE NEU1 LEVELS

Demonstrating a link between NEU1-downregulation and SARS-CoV-2 infectivity will allow not only to better understand aspects of pathogenesis of the COVID-19 disease, but also to test unconventional therapies aimed to raise NEU1 activity in patients. The need for these types of therapeutic approaches is dictated by the lack of any target therapy that could directly restore or increase NEU1 levels.<sup>49</sup> We have recently demonstrated that dietary supplements, like betaine, a natural amino acid derivative, as well as inhibitors of histone deacetylases,<sup>73,74</sup> increased the levels of mutant NEU1 when added to sialidosis fibroblasts or administered to mouse models of type I sialidosis with residual NEU1 activity.<sup>73,74</sup> Thus, it is conceivable that, if a correlation is found between NEU1 levels and extent and severity of viral infection, unconventional dietary compounds combined with conventional therapeutic approaches may prove beneficial for modulating NEU1 levels and ultimately for fighting infectious diseases like COVID-19.

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## AUTHOR CONTRIBUTIONS

A. Bongiovanni and A. d'Azzo conceptualized the hypothesis. A. Bongiovanni, A. Cusimano, I. Annunziata, and A. d'Azzo wrote the paper.

## ORCID

Alessandra d'Azzo  <https://orcid.org/0000-0003-1747-7502>



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