

University of Groningen

## Can ACE2 expression explain SARS-CoV-2 infection of the respiratory epithelia in COVID-19?

Nawijn, Martijn C; Timens, Wim

*Published in:*  
Molecular Systems Biology

*DOI:*  
[10.15252/msb.20209841](https://doi.org/10.15252/msb.20209841)

**IMPORTANT NOTE: You are advised to consult the publisher's version (publisher's PDF) if you wish to cite from it. Please check the document version below.**

*Document Version*  
Publisher's PDF, also known as Version of record

*Publication date:*  
2020

[Link to publication in University of Groningen/UMCG research database](#)

*Citation for published version (APA):*

Nawijn, M. C., & Timens, W. (2020). Can ACE2 expression explain SARS-CoV-2 infection of the respiratory epithelia in COVID-19? *Molecular Systems Biology*, 16(7), [e9841]. <https://doi.org/10.15252/msb.20209841>

### Copyright

Other than for strictly personal use, it is not permitted to download or to forward/distribute the text or part of it without the consent of the author(s) and/or copyright holder(s), unless the work is under an open content license (like Creative Commons).

### Take-down policy

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

*Downloaded from the University of Groningen/UMCG research database (Pure): <http://www.rug.nl/research/portal>. For technical reasons the number of authors shown on this cover page is limited to 10 maximum.*

# Can ACE2 expression explain SARS-CoV-2 infection of the respiratory epithelia in COVID-19?

Martijn C Nawijn\*  & Wim Timens

**Infection with severe acute respiratory syndrome coronavirus-2 (SARS-CoV-2) leads to coronavirus disease 2019 (COVID-19), which poses an unprecedented worldwide health crisis, and has been declared a pandemic by the World Health Organization (WHO) on March 11, 2020. The angiotensin converting enzyme 2 (ACE2) has been suggested to be the key protein used by SARS-CoV-2 for host cell entry. In their recent work, Lindskog and colleagues (Hikmet *et al*, 2020) report that ACE2 is expressed at very low protein levels—if at all—in respiratory epithelial cells. Severe COVID-19, however, is characterized by acute respiratory distress syndrome and extensive damage to the alveoli in the lung parenchyma. Then, what is the role of the airway epithelium in the early stages of COVID-19, and which cells need to be studied to characterize the biological mechanisms responsible for the progression to severe disease after initial infection by the novel coronavirus?**

*Mol Syst Biol.* (2020) 16: e9841

See also: F Hikmet *et al*

SARS-CoV-2 uses angiotensin-converting enzyme 2 (ACE2) as receptor for docking and cell entry via the spike (S) glycoprotein. To elucidate the pathogenesis of COVID-19, it is of critical importance to identify the cells infected by SARS-CoV-2, as well as the mechanisms of viral entry, replication, and the ensuing cellular damage leading to the induction of innate and adaptive antiviral responses. Several recent studies report on the expression of ACE2 mRNA

in tissues or individual cells, identifying the upper airway epithelium as the most likely site of entry for SARS-CoV-2 (Lukassen *et al*, 2020; Sungnak *et al*, 2020). In their recent study, Hikmet *et al* (2020) present a much-needed updated systematic evaluation of ACE2 expression in a large range of tissues, at the protein as well as the RNA level, confirming the expression in a number of epithelial barrier tissues. Importantly, the authors show that ACE2 expression in the respiratory tract is very limited compared to other barrier tissues, and reproducible staining is observed in nasal epithelial cells with only one of the two antibodies used in this study, prompting the authors to claim that none or only very low levels of ACE2 protein is present in the normal respiratory system.

Identification of the entry site of SARS-CoV-2 in COVID-19 patients will facilitate focused research efforts to test whether and how the early response to viral infection governs progression to severe and fatal disease. Such insight into the pathogenesis of early stages of COVID-19 might aid the identification of biomarkers for those patients most likely to progress to severe disease, and guide design of therapeutic interventions aiming at preventing severe disease in its variations. Finally, it might offer opportunities to further improve personal protection measures for those (healthcare) professionals at risk of attracting the virus. Several reports have started to explore the response of airway epithelial cells to SARS-CoV-2 infection. For instance, recent data indicate that primary airway epithelial cells have a relatively modest type-1 interferon response to SARS-CoV-2

infection (Blanco-Melo *et al*, 2020), and their interactions with the immune system correlate to disease severity (Chua *et al*, 2020). However, do we need to re-evaluate a role for the airway epithelium as primary site of infection with SARS-CoV-2 based on the findings reported by Hikmet *et al*?

Their study reports on RNA and protein expression of ACE2 in a range of human tissues based on the Human Protein Atlas (HPA) resource. The tissue-level RNA expression data align well with the single-cell RNA sequencing (scRNA-Seq) datasets re-analyzed by the authors, although the nasal epithelium and the conjunctiva of the eye are lacking from the HPA dataset while showing relatively high ACE2 expression in the single-cell analyses (Sungnak *et al*, 2020). The protein expression data for ACE2, however, represent the most relevant part of this study, with contrasting data in literature for ACE2 expression in the respiratory system (Hamming *et al*, 2004; Jia *et al*, 2005). The authors convincingly show expression in a range of tissues including duodenum, small and large intestine, the proximal tubule cells of the kidney, glandular cells in the gallbladder, and cardiomyocytes, as well as the conjunctiva of the eye. In the normal respiratory system, however, the authors observe positive staining only with one of two antibodies used, in ciliated cells in the nasal cavity (in all subjects tested), in the bronchus (in a subset of the subjects tested), and in a subset of alveolar type-2 epithelial cells, and conclude that ACE2 protein expression in lung is very limited at best.

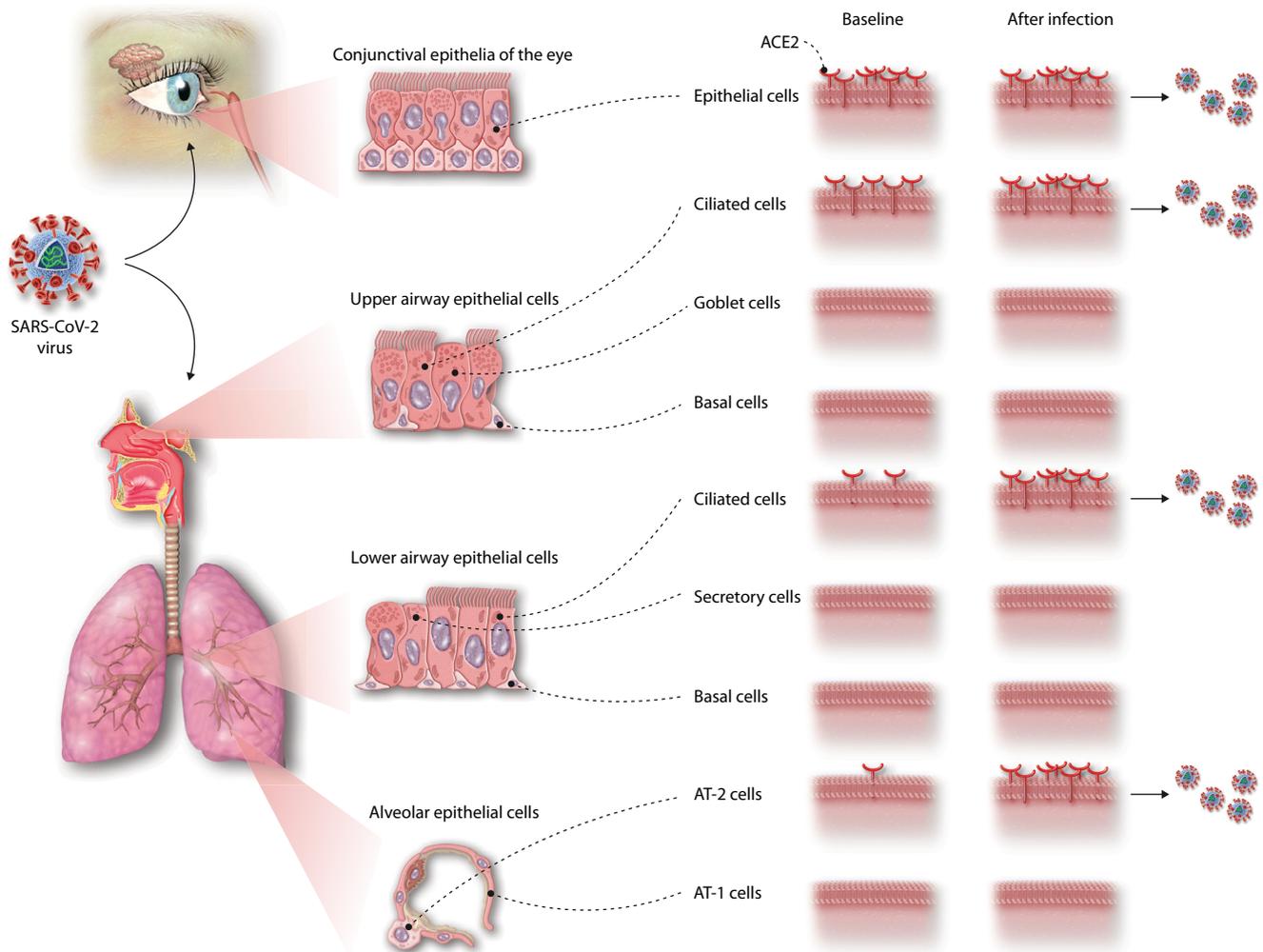
The baseline for ACE2 protein expression in normal tissue as reported by Hikmet *et al*

is important for unraveling the pathogenesis of COVID-19. Is the reported low level of ACE2 protein expression sufficient to support a role for the (upper) airway epithelium, in combination with the conjunctiva of the eye, as a first site of infection by SARS-CoV-2 as a prelude to developing COVID-19? The nasal epithelial staining with the more sensitive antibody is reproducible, and the staining pattern matches that observed in the other epithelial barrier tissues. For the lower airways, however, staining with this antibody is not reproducible between donors. The antibody used in the earlier studies reporting positive staining in lower airways (Hamming *et al*, 2004) was raised against a peptide sequence partly overlapping with the peptide used to generate the

antibodies Hikmet *et al* used. Differences in recognition (and consequently in level of immunohistological expression) can also be due to differences in folding of the protein and thus differences in epitope expression. Moreover, ACE2 has been shown to be an interferon-induced gene (Ziegler *et al*, 2020), and ACE2 levels in airway epithelial cells dramatically increase during SARS-CoV-2 infection (Chua *et al*, 2020). The presence of viral RNA has been observed in ciliated but not in secretory epithelial cells in the airways of COVID-19 patients (Hou *et al*, 2020), which is in line with the ACE2 staining pattern reported in this manuscript (Hikmet *et al*, 2020), indicating that ACE2 expression might be sufficient to allow infection of the ciliated airway epithelial cells by

SARS-CoV-2. In such a scenario (Fig 1), low-level protein expression in upper airway epithelial cells facilitates infection of ciliated cells, followed by a rapid interferon-induced increase of ACE2 expression in lower airways and lung parenchyma that then sets the stage for viral spread across the respiratory mucosa and which could contribute to severe COVID-19.

In addition to baseline expression and changes thereof induced by SARS-CoV-2 infection, the effects of external factors and pre-existing lung disease on ACE2 expression need to be considered. For instance, smoking has been shown to increase ACE2 expression in the respiratory tract (Smith *et al*, 2020). Expression of ACE2 in patients with pre-existing lung disease also needs to



**Figure 1. Proposed events during SARS-CoV-2 infection of the respiratory tract.** (A) At baseline, ACE2 expression is present in the conjunctival epithelia of the eye and at low levels in the ciliated epithelial cells in the upper airways, allowing SARS-CoV-2 infection at these initial sites. (B) The antiviral response induced upon SARS-CoV-2 infection induces a marked upregulation of ACE2 expression, allowing the SARS-CoV-2 to spread across the respiratory mucosa and into the parenchyma of the lung, where it can infect type-2 alveolar epithelial cells.

be systematically evaluated, as very often these disorders are accompanied by significant increases in alveolar type-2 epithelial cells, that are in the center of the main COVID-19 lung pathology. Therefore, further studies are urgently needed on the role of respiratory epithelial cells during infection with SARS-CoV-2 and the early stages of COVID-19, in particular in risk groups and patients with pre-existing disease, to identify the complex mechanisms that lead to the exaggerated immune response seen in severe disease.

## References

- Blanco-Melo D, Nilsson-Payant BE, Liu W-C, Uhl S, Hoagland D, Møller R, Jordan TX, Oishi K, Panis M, Sachs D *et al* (2020) Imbalanced host response to SARS-CoV-2 drives development of COVID-19. *Cell* 181: 1036–1045.e9
- Chua RL, Lukassen S, Trump S, Hennig BP, Wendisch D, Pott F, Debnath O, Thürmann L, Kurth F, Völker MT *et al* (2020) COVID-19 severity correlates with airway epithelium-immune cell interactions identified by single-cell analysis. *Nat Biotechnol* <https://doi.org/10.1038/s41587-020-0602-4>
- Hamming I, Timens W, Bulthuis MLC, Lely AT, Navis GJ, van Goor H (2004) Tissue distribution of ACE2 protein, the functional receptor for SARS coronavirus. A first step in understanding SARS pathogenesis. *J Pathol* 203: 631–637
- Hikmet F, Méar L, Edvinsson A, Micke P, Uhlén M, Lindskog C (2020) The protein expression profile of ACE2 in human tissues. *Mol Syst Biol* 16: e9610
- Hou YJ, Okuda K, Edwards CE, Martinez DR, Asakura T, Dinnon KH III, Kato T, Lee RE, Yount BL, Mascenik TM *et al* (2020) SARS-CoV-2 reverse genetics reveals a variable infection gradient in the respiratory tract. *Cell* <https://doi.org/10.1016/j.cell.2020.05.042>
- Jia HP, Look DC, Shi L, Hickey M, Pewe L, Netland J, Farzan M, Wohlford-Lenane C, Perlman S, McCray PB (2005) ACE2 receptor expression and severe acute respiratory syndrome coronavirus infection depend on differentiation of human airway epithelia. *J Virol* 79: 14614–14621
- Lukassen S, Chua RL, Trefzer T, Kahn NC, Schneider MA, Muley T, Winter H, Meister M, Veith C, Boots AW *et al* (2020) SARS-CoV-2 receptor ACE2 and TMPRSS2 are primarily expressed in bronchial transient secretory cells. *EMBO J* 39: e105114
- Smith JC, Sausville EL, Girish V, Yuan ML, Vasudevan A, John KM, Sheltzer JM (2020) Cigarette smoke exposure and inflammatory signaling increase the expression of the SARS-CoV-2 receptor ACE2 in the respiratory tract. *Dev Cell* 53: 514–529.e3
- Sungnak W, Huang N, Bécavin C, Berg M, Queen R, Litvinukova M, Talavera-López C, Maatz H, Reichart D, Sampaziotis F *et al* (2020) SARS-CoV-2 entry factors are highly expressed in nasal epithelial cells together with innate immune genes. *Nat Med* 26: 681–687
- Ziegler CGK, Allon SJ, Nyquist SK, Mbano IM, Miao VN, Tzouanas CN, Cao Y, Yousif AS, Bals J, Hauser BM *et al* (2020) SARS-CoV-2 receptor ACE2 is an interferon-stimulated gene in human airway epithelial cells and is detected in specific cell subsets across tissues. *Cell* 181: 1016–1035.e19



**License:** This is an open access article under the terms of the Creative Commons Attribution 4.0 License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited.