

# **SARS-CoV-2 infection demonstrated in a human lung bronchioalveolar tissue model**

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Development of an in vitro human-derived tissue model for studying virus infection and disease progression in the alveolar cells of the lungs responsible for oxygen and carbon dioxide exchange with the blood

might enable the study of possible therapies for acute respiratory distress syndrome (ARDS) triggered by SARS-CoV-2. Researchers in the Netherlands have demonstrated that the SARS-CoV-2 replicates efficiently in their model resembling the human bronchioalveolar system that is thought to play a critical role in progression of infection towards pneumonia and ARDS.

It is already established that in people infected with COVID-19 or some other respiratory viruses, alveolar injury can trigger a cascade of events that leads to ARDS, restricting transport of oxygen into the blood to dangerously low levels. There is also mounting evidence that the epithelium lining the alveoli plays a major role in progression of COVID-19. However, in vitro models for replicating [disease progression](#) in the alveoli of human lungs have proven difficult to establish, especially models that are also permissive to SARS-CoV-2 infection. This has greatly limited our understanding of COVID-19.

The Dutch team has now remedied this deficiency through application of self-renewing organoid models containing stem [cells](#) capable of differentiating into relevant cell types for study of disease processes. Organoids are tiny 3-D tissues typically around 2 mm in diameter across derived from stem cells to mirror the complex structures of an organ, or at least to express selected aspects of it to meet a given biomedical research objective. Such organoids can then provide continuous sources of 2-D tissues that mimic more accurately the geometry or cellular alignment of the structures under study.

A self-renewing organoid [model](#) for the epithelium of the airways conducting the gases, has already been developed by the same team, but the alveolar epithelium has proven a greater challenge to generate so far. The Dutch team has overcome this challenge and developed a 2-D 'air interface' system comprising a basal layer of [stem cells](#) in contact with the culture media and a top layer exposed to the air just as it would

be in the lungs.

Multiple cultures were generated and infected successfully by SARS-CoV-2 targeting primarily alveolar type-II-like cells, known as ATII-L, confirmed by Transmission Electron Microscopy (TEM), surface marker stainings and single-cell sequencing. The study then shed light on the sequence of events following infection.

The study also identified through messenger RNA expression analysis a cellular immune response to the virus by infected cells. When the cultures were treated with the antiviral signaling molecule interferon lambda early in infection, SARS-CoV-2 replication was almost completely blocked, indicating that—when timed right—interferon lambda could be an effective treatment. These results also indicate that these cultures could be helpful for the development of a therapeutic intervention against [acute respiratory distress syndrome](#) (ARDS) from COVID-19.

**More information:** Mart M. Lamers et al, An Organoid-derived Bronchioalveolar Model for SARS-CoV-2 Infection of Human Alveolar-type II-like Cells, *The EMBO Journal* (2020). [DOI: 10.15252/embj.2020105912](https://doi.org/10.15252/embj.2020105912)

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