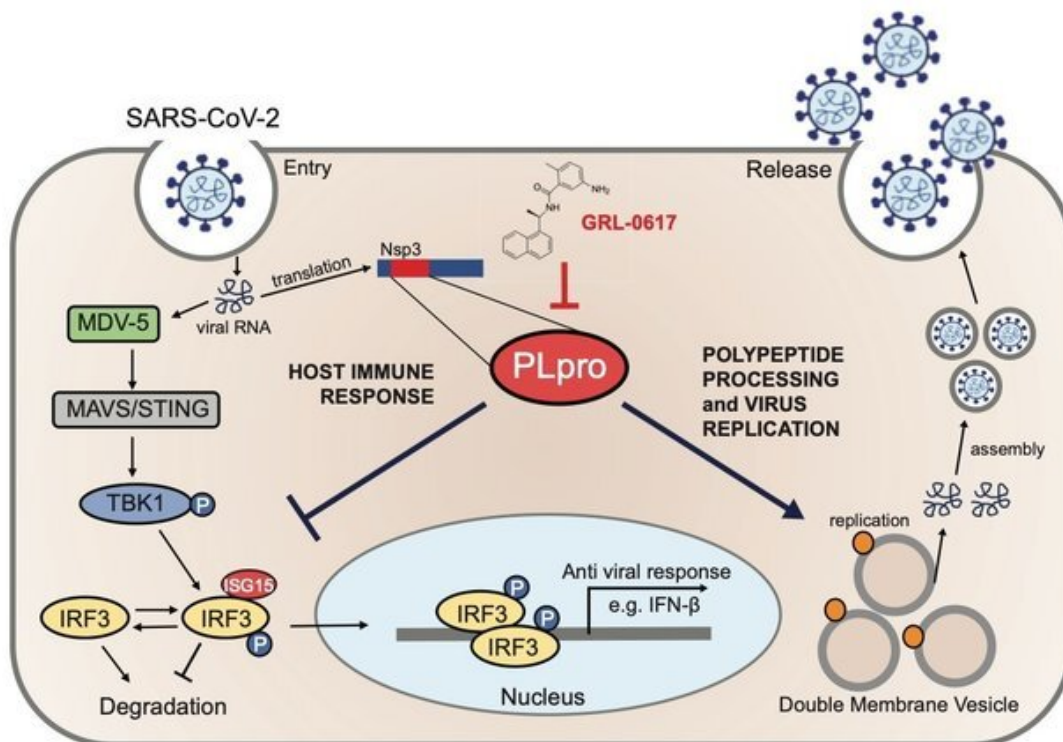


COVID-19: Anti-viral strategy with double effect

July 29 2020, by Brigitte Holfelder



The protease PLpro of Sars-CoV-2 plays a central role in the viral life cycle. By cleaving also proteins of infected cells, PLpro interferes with the innate immune response of the host. Inhibition of PLpro fosters the anti-viral interferon response and reduces viral replication in infected cells. Credit: Donghyuk Shin, Max Planck Fellow, MPI of Biophysics, Nature 2020

When the SARS-CoV-2 virus penetrates human cells, it lets the human host cell produce proteins for it. One of these viral proteins, called PLpro, is essential for the replication and rapid spread of the virus. An international team of researchers led by Goethe University, University Hospital Frankfurt and Max Planck Institute of Biophysics has now discovered that the pharmacological inhibition of this viral enzyme not only blocks virus replication but also strengthens the anti-viral immune response at the same time.

In the case of an infection, the SARS-CoV-2 [virus](#) must overcome various defense mechanisms of the human body, including its non-specific or innate immune defense. During this process, infected body [cells](#) release messenger substances known as type 1 interferons. These attract natural killer cells, which kill the infected cells.

One of the reasons the SARS-CoV-2 virus is so successful—and thus dangerous—is that it can suppress the non-specific immune response. In addition, it lets the human cell produce the viral protein PLpro (papain-like protease). PLpro has two functions: It plays a role in the maturation and release of new viral particles, and it suppresses the development of type 1 interferons. The German and Dutch researchers have now been able to monitor these processes in cell culture experiments. Moreover, if they blocked PLpro, virus production was inhibited and the innate immune response of the [human cells](#) was strengthened at the same time. The molecular dynamics calculations of Laura Schulz and Ahmad Reza Mehdipour at the Max Planck Institute for Biophysics and the Max Planck Computing and Data Facility in Garching have made a major contribution to elucidating the underlying mechanism and better understanding the binding of the inhibitor.

Promising therapeutic strategy

Ivan Đikić explains: "We used the compound GRL-0617, a non-covalent

inhibitor of PLpro, and examined its mode of action very closely in terms of biochemistry, structure and function. We concluded that inhibiting PLpro is a very promising double-hit therapeutic strategy against COVID-19. The further development of PLpro-inhibiting substance classes for use in [clinical trials](#) is now a key challenge for this therapeutic approach."

Another important finding from this work is that the viral protein PLpro of SARS-CoV-2 cleaves off ISG-15 (interferon-stimulated gene 15) from cellular proteins with a higher level of activity than the SARS equivalent, which leads to greater inhibition of type I interferon production. This is concordant with recent clinical observations which show that COVID-19 exhibits a reduced interferon response in comparison to other respiratory viruses such as influenza and SARS.

To understand in detail how inhibiting PLpro stops the virus, researchers in Frankfurt, Munich, Mainz, Freiburg and Leiden have worked closely together and pooled their biochemical, structural, IT and virological expertise.

Sandra Ciesek, Director of the Institute of Medical Virology at University Hospital Frankfurt, explains that the papain-like protease is an extremely attractive anti-viral goal for her as a physician because its inhibition would be a "double strike" against SARS-CoV-2. She highlights the excellent collaboration between the institutes: "Especially when investigating a new clinical picture, everyone profits from interdisciplinary collaboration as well as different experiences and viewpoints." Gerhard Hummer from the Max Planck Institute for Biophysics adds: "This work is the result of a broad-based collaboration. Under [difficult circumstances](#) at the peak of the first COVID-19 wave, many researchers here discovered how the virus interferes with our immune response and at the same time identified possible therapeutic approaches in the form of an inhibitor."

More information: Donghyuk Shin et al. Inhibition of papain-like protease PLpro blocks SARS-CoV-2 spread and promotes anti-viral immunity, (2020). [DOI: 10.21203/rs.3.rs-27134/v1](https://doi.org/10.21203/rs.3.rs-27134/v1)

Donghyuk Shin et al, Papain-like protease regulates SARS-CoV-2 viral spread and innate immunity, *Nature* (2020). DOI: 10.1038/s41586-020-2601-5

Provided by Max Planck Society

Citation: COVID-19: Anti-viral strategy with double effect (2020, July 29) retrieved 26 April 2024 from <https://medicalxpress.com/news/2020-07-covid-anti-viral-strategy-effect.html>

This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.