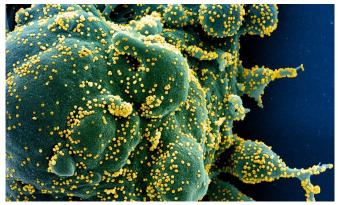


Clinical histone deacetylase inhibitors are effective against COVID-19

14 December 2020, by Li Yuan



ACE2 receptor by its spike protein during viral infection, such alternations inhibited the ACE2-S protein binding and prevented host cell entry of SARS-CoV-2.

Inspired by this result, the team then screened 18 commercially available HDAC inhibitors and studied their efficacy in inhibiting the entry of SARS-CoV-2 into cells. They found that four inhibitors, i.e., panobinostat, givinostat hydrochloride monohydrate, CAY10603 and sirtinol are noticeably effective.

Colorized scanning electron micrograph of an apoptotic cell (green) heavily infected with SARS-COV-2 virus particles (yellow), isolated from a patient sample. Image captured at the NIAID Integrated Research Facility (IRF) in Fort Detrick, Maryland. Credit: NIH/NIAID

The lowest half maximal effective concentration (EC50) of viral transduction inhibition achieved by these four HDAC drugs is about 3 ?M, which is far less than EC50 obtained from other non-effective drugs (all more than 100?M).

COVID-19, caused by the SARS-CoV-2 virus, has posed serious health threats to humanity and jeopardized the global economy.

The possible mechanism of these HDAC inhibitors that makes them effective against SARS-CoV-2 was revealed by the cellular signaling network analysis.

However, no effective drugs are available to treat COVID-19 currently and there is a great demand to fight against it.

This work was published as a cover story in ACS Pharmacology & Translational Science.

By combining computational screening and an efficient cellular pseudotyped virus system, a research team led by Prof. YUAN Shuguang from the Shenzhen Institutes of Advanced Technology (SIAT) of the Chinese Academy of Sciences has confirmed that clinical histone deacetylase (HDAC) inhibitors can efficiently prevent SARS-CoV-2 and potentially be used to fight against COVID-19.

More information: Ke Liu et al. Clinical HDAC Inhibitors Are Effective Drugs to Prevent the Entry of SARS-CoV2, ACS Pharmacology & Translational Science (2020). DOI: 10.1021/acsptsci.0c00163

The researchers found that several HDAC inhibitors could bind to human angiotensin I converting enzyme 2 (ACE2) on the <u>cell surface</u>, which in turn resulted in overall structural changes of ACE2. Since SARS-CoV-2 recognizes human

Provided by Chinese Academy of Sciences



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