COVID-19 (COronaVIrus disease 2019) background:

First case identified in December 2019 in Wuhan China

caused by SARS-CoV-2 (severe acute respiratory syndrome coronavirus 2) virus

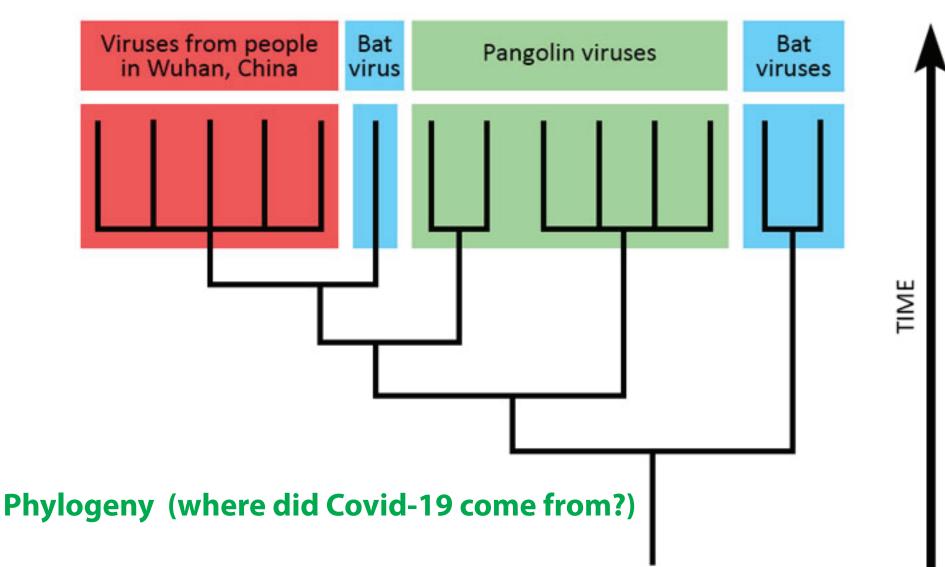
SARS-CoV-2 virus is a positive-sense single-stranded RNA virus

"positive-sense" RNA virus -- can act as mRNA and therefore can be **directly** translated into protein

"negative-sense" RNA virus has genome being complementary to mRNA so that mRNA must be synthesized via RNA-dependent RNA polymerase

SARS-CoV-2 virus: 29,903 bases (Large for RNA virus!) Human genome: about 3.2 billion bases

CORONAVIRUS EVOLUTIONARY TREE



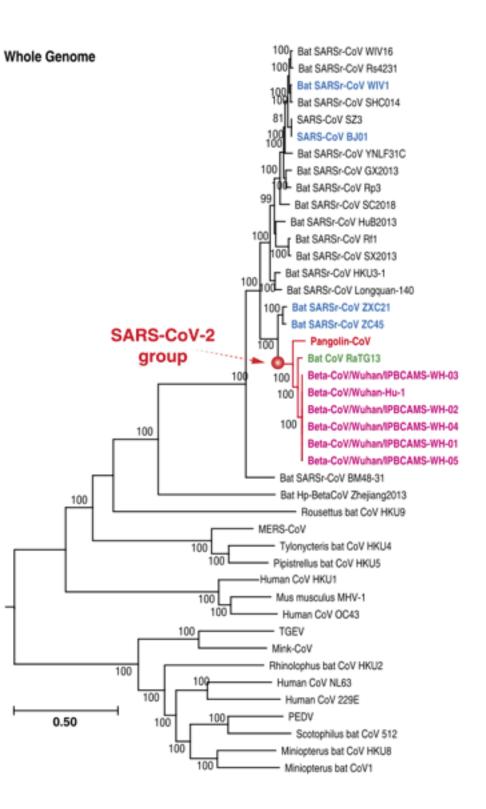
Zoonotic Transfer History

image from https://evolution.berkeley.edu/evolibrary/news/200305_coronavirus

Phylogeny of the SARS-like coronaviruses. From: Zhang et al. (Current Biology, 2020) [Redirecting] (https://doi.org/10.1016/j.cub.2020.03.022) [copied in turn from Dr. David Rasmussen]

Update: An "in press" Nature article Cave bats in Laos found to harbor closer virus sequences to human pandemic sequences than those previously known. Also, different regions of viral genome of that causing human pandemic seem to have come from different ancestors ("a mosaic history").

(Temmam, S. et al. Bat coronaviruses related to SARS-CoV-2 and infectious for human cells. Nature https:// doi.org/10.1038/s41586-022-04532-4. 2022)



Is Covid-19 genetically engineered? Andersen et al. 2020. The Proximal Origin of SARS-CoV2. Nature Medicine.

RaTG13 viral sequence from a bat is 96% identical to HCoV-19 genome.

But (!), virus from pangolins is more similar to Receptor Binding Domain (RBD) in HCoV-19 & 6 key RBD residues from pangolin viruses identical to HCoV-19 ones (only 1 of 6 identical to HCoV-19 in closest bat sequence).

RBD binds to human ACE receptor to allow virus to enter cells.

HCoV-19 RBD differs from previously identified optimal RBD sequence

(Also, HCoV-19 genome substantially different than coronavirus genomes studied by molecular geneticists)