

Chloroquine and COVID-19: A Light at the End of the Tunnel, or is it Another Train?

Editorial

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ARTICLE INFO	ABSTRACT
Received: 29 Mar. 2020	Over the last a few decades, the world has faced several viral disease outbreaks including Hendra, Nipah, sever
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Over the last a few decades, the world has faced several viral disease outbreaks including Hendra, Nipah, severe acute respiratory syndrome (SARS), Influenza (H1N1), Middle East respiratory syndrome (MERS), Zika and most recently coronavirus disease 2019 (COVID-19). The economical and health impacts of the outbreaks were limited to certain affected areas only, but the recent outbreak of COVID-19 seems to have a global impact on human health, health resources and global economy.

COVID-19 is a member of the species of the coronavirus family that cause *severe acute respiratory syndrome-related diseases*. COVID-19 is a zoonotic virus that belongs to the Coronavirus family, which belongs to the *Coronaviridae* suborder (1). Currently, there is no vaccine or treatment for COVID-19. The high infectivity and lethality of COVID-19 necessitated its classification as a biological safety level 4 (BSL-4) pathogen. Since its emergence in late 2019 in Wuhan-China, COVID-19 infected approximately 678,905 including 31,771 deaths worldwide (as of March 29 2020) (2).

There is an extensive effort across the globe to develop novel coronavirus vaccine or antiviral treatment to tackle the spread of the mysterious and previously unknown virus infection. While we remain hopeful that the current effort will be fruitful, we speculate that at best an effective treatment may not be available before yearend. Nonetheless, one of the most promising drug candidates is chloroquine, an FDA approved therapy for the treatment and prevention of malaria infection, which has received an international attention for its potential effect as an antiviral against COVID-19. The Chinese National Centre for Biotechnology Development announced that the results of 100 patients treated with chloroguine showed improved CT scan of lung, reduction in fever and a shorter recovery time compared with other groups (3). Also, a study from France that used the drug in a small number of infected patients claimed to have positive clinical outcomes (4).

The use of chloroguine as an antiviral is not an entirely new phenomenal. Almost four decades ago, Coombs and colleagues claimed that the use of chloroquine significantly reduced viral yields of Sindbis virus in BHK cells (5). Similarly, the drug was tested in vitro against SARS virus by Vincent et al., 2005, which claimed that chloroquine shows potent antiviral activity against SARS virus (6). The authors claimed that the antiviral activity of chloroquine is partly due to its effect in increasing endosomal pH and its ability to glycosylate the viral entry receptor, angiotensin converting enzyme. Also, in 2013 Yan et al, reported that chloroquine is highly effective in treating avian influenza A H5N1 virus in animals (7). Furthermore, in collaboration with Weil Cornell Medical College, we showed that chloroquine was able to inhibit Nipah and Hendra virus infections in vitro (8). However, the use of chloroquine against Nipah infected ferrets, did not prevent the disease (8).

In conclusion, despite the fact that the drug has been in use for more than half a century and was shown in vitro to have potent antiviral activity against several viruses including COVID-19, chloroquine failed to treat several viral infections in animal and human studies. Therefore, in the absence of randomized control clinical trials, we should be careful not to draw early conclusions, as the light we see may turnout to be another train.

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