## Letter to Editor

## What is the Role of the Anti-Parasitic Drugs in the Treatment of Coronavirus Disease 2019?

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## **Dear Editor-in-chief**

Severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) is a rapidly emerging viral infection causing coronavirus disease 2019 (COVID-19). The first report of the novel coronavirus, SARS-CoV-2, originated in Wuhan, China, in early December 2019<sup>1</sup>. Currently, there is no U.S. food and drug administration (FDA) approved drugs for the treatment of patients with COVID-19. A great deal of effort is ongoing to find effective therapeutics and preventive measures against this transmissible virus with high mortality. As result, available data are limited, and there are minimal randomized controlled trial (RCT) data on the efficacy of antiviral or immunomodulatory agents for the treatment of COVID-19<sup>2</sup>. Most of the treatment protocols are based on observational studies and anecdotic reports<sup>3,4</sup>.

Until now, there is no specific drug therapy available for the treatment of COVID-19. However, some potential anti-parasitic drugs like hydroxychloroquine and azithromycin, antifilarial drug Ivermectin and antiviral drugs have been tested by many research groups worldwide for their possible effect against the COVID-19<sup>5</sup>. Ivermectin was a revolutionary drug in the 1980s, the forerunner of a new group of antiparasitic agents with activity against both parasitic nematodes and arthropods. Ivermectin has valuable public health applications for controlling strongyloidiasis and scabies and filariasis, through its effect on transmission<sup>6,7</sup>. Recent reports suggested that the treatment of COVID-19 by the single dose of Ivermectin was found to reduce the viral load up to 5000 fold in vitro culture within 48 h<sup>8</sup>. Mechanism of action of this drug against COVID-19 is still unknown, but the impotant features of this drug is that no toxicity was observed during in vitro culture.

Chloroquine (CQ) and hydroxychloroquine (HCQ) have been used to treat malaria for 70 years. Chloroquine (CQ) has showed low toxicity with strong inhibition on MERS-CoV and SARS-CoV<sup>9</sup>. Chloroquine likely accumulates in lysosomes, where it sequesters protons and increases the pH. The drug modulates immune response by interacting variety of host proteins and cellular processes.

The HCQ is derivative of CQ, and is the first category drug which is prescribes for COVID-19 infection therapy<sup>10</sup>. This drug is a first line drug for malaria treatment and also has been used for rheumatoid arthritis and systemic lupus erythematosus treatment<sup>11</sup>. HCQ specific feature is affecting acid balance and can inhibit many enzymes. Moreover, it can obstruct the viral post-translational modifications and glycosyl-transferases<sup>12</sup>.

Many Research centers and institutes all around the world have announced institutional guidelines for offlabel use of drugs for COVID-19, including CQ and HCQ with different dosages and duration for either treatment or prophylaxis, but there is no standard recommendation for prescribing these medications for this disease. In addition, CQ and HCQ have many side effects, including cardiac toxicity (QT prolongation, torsade de pointes, and ventricular arrhythmia), which may be harmful in the elderly, who are also high risk for COVID-19<sup>13</sup>.

However, it is still not clear that whether these drugs have a better therapeutic effect, when compared to other drugs or combination therapy of multiple drugs. Consequently, this editorial makes comprehensive view on anti-parasitic drugs, which can be used in the current treatment COVID-19 disease.

**Keywords:** COVID-19, Treatment, Anti-parasitic drugs, Hydroxychloroquine, Chloroquine

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