LETTER TO EDITOR

Dear Editor,

Chloroquine (CQ) is considered as a drug that has been significantly marketed versus malaria and observed in 2006 that this drug has an antimicrobial potential for the mechanism and chloroquine function increases the pH of the endosomal [1]. With a higher pH of what is required for virus/cell fusion, it blocks the infection and prevents the progression of the disease. This is due to the SARS-COV-2 cellular receptor glycoproteins that are connected to their items [2]. But the hydroxychloroquine can inhibit the stage before the cellular binding in the cell surface receptor, in a study in a study by a group of Chinese researchers and scientists about the efficiency and function of chloroquine Laboratory studies have been carried out using VERO E6 cells infected with SARS-COV-2 in polyethylene infection (MOI) 0.05, chloroquine has been very effective in reducing the virus and has an impressive effect and can infect SARS-COV-2 block at low concentrations (effective concentration of maximum (EC50) 1.13μM and the concentration of semi-cytotoxicity (CC50) larger than 100μM). Another article showed that chloroquine could prevent the proliferation of HCOV-229E in laboratory conditions in the culture of lung epithelial cells and can also be effective against the Coronavirus of the Middle East Respiratory Syndrome (Mers-Cov) in laboratory conditions. Hydroxychloroquine (HCQ) with a very similar chemical structure with chloroquine is one of the moderating diseases of anti-rheumatic drugs used to treat many [3]. The use of this lopinavir and ravidar, which actually produced to deal with the AIDS virus, has also not affected the status of citizens with corona. The experiments showed that these drugs, which are usually sold by Kaltra, have reduced the mortality rate caused by corona, rather than the risk of sub-oxygen. This treatment method has not even shorter patient’s hospitalization. In another study, anti-HIV drugs, lopinavir and paint, may have a therapeutic effect on Covid-19 disease [3]. This means that the therapeutic effect of Ritonavir and Lopinavir on Covid-19 may be largely due to its inhibitory effect on CEP.C30 of the corona virus, while Ritonavir may have more ability and performance, the inhibitory effect of Darney on SARS-COV-2 and the effect of its potential therapeutic therapy may be largely due to its inhibitory effect on PLVP (peptic viral protease) [1]. Lopinavir is metabolized by cytochrome P4503A (CYP3A) in the liver. This drug is always used with Ritonavir to reduce the dose of lopinavir and increase the level of lopinavir in plasma because Ritonavir prevents the Isoenzyme CYP 3A. Lopinavir and Ritonavir viral protease inhibitors have been selected as the second line for the treatment of HIV-1 infection in children and adults with lower side effects with lower side effects of steroids and hospital infections in patients initially treated with Lopinavir/Ritonavir. They have been observed, also, the results indicate a reduction in viral load and increase the number of peripheral lymphocytes. The findings of laboratory and clinical studies, along with the specifications of the availability and immunity of Lopinavir/Ritonavir and interferon beta 1b (IFN-β1b), show that the combination of these factors to treat patients with Mers-Cov has a potential effect. The question of whether the previous approach of Lopinavir-Ritonavir in Covid-19 can have clinical benefits, is an important issue that requires more study. A new report showed that the average duration of virus release in COVID-19 in patients with severe illness was 20 days and can also take up to 37 days. There is no proven evidence that Lopinavir-Ritonavir has a significant antiviral effect [4].

REFERENCES