No difference in biomarkers of ischemic heart injury and heart failure in patients with COVID-19 who received treatment with chloroquine phosphate and those who did not

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Abstract

Background
Chloroquine was promoted as a COVID-19 therapeutic early in the pandemic. Most countries have since discontinued the use of chloroquine due to lack of evidence of any benefit and the risk of severe adverse events. The primary aim of this study was to examine if administering chloroquine during COVID-19 imposed an increased risk of ischemic heart injury or heart failure.

Methods
Medical records, laboratory findings, and electrocardiograms of patients with COVID-19 who were treated with 500 mg chloroquine phosphate daily and controls not treated with chloroquine were reviewed retrospectively. Controls were matched in age and severity of disease.

Results
We included 20 patients receiving chloroquine (500 mg twice daily) for an average of five days, and 40 controls. The groups were comparable regarding demographics and biochemical analyses including C-reactive protein, thrombocytes, and creatinine. There were no statistically significant differences in cardiac biomarkers or in electrocardiograms. Median troponin T was 10.8 ng/L in the study group and 17.9 ng/L in the control group, whereas median NT-proBNP was 399 ng/L in patients receiving chloroquine and 349 ng/L in the controls.
Conclusions

We found no increased risk of ischemic heart injury or heart failure as a result of administering chloroquine. However, the use of chloroquine to treat COVID-19 outside of clinical trials is not recommended, considering the lack of evidence of its effectiveness, as well as the elevated risk of fatal arrhythmias.

1. Introduction

Coronavirus disease 2019 (COVID-19) is an infection caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), with a symptomatology that varies from no (or very mild) symptoms in the upper airways to severe viral pneumonia with respiratory failure. According to current understanding, approximately 80% of symptomatic cases are mild, 15% severe, and 5% critical [1], while the proportion of asymptomatic cases is estimated to be 30% to 40% [2]. The most common complications in patients with severe disease are interstitial pneumonia with respiratory failure, pulmonary thrombosis, arrhythmia, acute cardiac, and acute kidney injury [3, 4]. The mortality among patients in intensive care varies between different studies [4]; in Sweden the mortality among patients requiring invasive ventilation is approximately 20% [5].

As of January 2021, the only drugs approved by the European Medicines Agency (EMA) and US Food and Drug Administration (FDA) are the antiviral drug remdesivir, dexamethasone (EMA only), convalescent plasma (FDA only), the janus kinase inhibitor baricitinib in combination with remdesivir (FDA only), and monoclonal antibodies (FDA only) [6, 7]. Multiple trials ongoing worldwide include new treatment options and drugs that have already been developed for other diseases [8, 9]. One of the drugs that has been widely administered is the antimalarial drug chloroquine, either as chloroquine phosphate or as the less toxic derivative hydroxychloroquine [9]. Because of the immunomodulatory effects of chloroquine, these drugs have long been used to treat inflammatory diseases such as rheumatoid arthritis and systemic lupus erythematosus. Chloroquine has been shown to affect cytokine production [10, 11], which raised hopes that it might mitigate the cytokine release syndrome believed to be associated with severe COVID-19 [12]. Studies have also shown a broad antiviral effect in vitro against several coronaviruses including SARS-CoV-2 [13]. The first positive reports of the clinical efficacy of chloroquine came from China in February 2020 [14], and shortly thereafter a small French study showed a positive effect of chloroquine on symptoms and viral clearance [15]. Since then, chloroquine has been used to a varying degree in many countries, including Sweden. The Department of Infectious Diseases at Sahlgrenska University Hospital, Gothenburg, discontinued the use of chloroquine phosphate at the end of March 2020 due to a lack of evidence regarding efficacy and the risk of severe adverse events. Several studies have since reported severe adverse events arising from the use of chloroquine, with or without the co-administration of azithromycin, to treat COVID-19, especially arrhythmias and QT-prolongation [16, 17]. While the potential benefits of chloroquine/hydroxychloroquine continue to be widely debated, after promising initial reports several clinical trials have been published with no positive results regarding viral clearance and clinical status as compared to standard of care [18]. The hydroxychloroquine arm of the Solidarity Trial, launched by WHO and partners, was permanently halted in June 2020 when data showed that, in comparison to standard of care, hydroxychloroquine did not result in a reduction of in-hospital mortality [19, 20].
To our knowledge, no studies of markers of cardiac injury during chloroquine treatment for COVID-19 have been published to date. The primary aim of our study was to examine whether patients with COVID-19 undergoing treatment with chloroquine phosphate had an increased risk of cardiotoxicity with regard to ischemic heart injury, heart failure, and arrhythmias, when compared to patients receiving standard treatment.

2. Methods

2.1 Study design and population

This is a retrospective study of patients with COVID-19 who received treatment with chloroquine phosphate (500 mg twice daily) and matched controls from the same time period who did not receive chloroquine phosphate. Inclusion criteria were a) confirmed diagnosis of COVID-19 by positive polymerase chain-reaction test (PCR) from the upper respiratory tract for SARS-CoV-2; b) admission to the Department for Infectious Diseases at Sahlgrenska University Hospital, Gothenburg, Sweden; and c) having received a minimum of two doses of chloroquine phosphate. Exclusion criteria were lack of testing for cardiac biomarkers (cardiac troponin T [cTnT] and N-terminal pro b-type natriuretic peptide [NT-proBNP]), or lack of stored blood samples from either the treatment period or the first four weeks of symptomatic infection. Controls were chosen from the same patient cohort, with chloroquine phosphate treatment as the exclusion criteria. Participants were matched as closely as possible for severity of disease and age.

2.2 Data collection and variables of the study

We collected data regarding sex, age, weight, height, comorbidities, medication, date of symptom onset, severity of disease, electrocardiogram (ECG) results, laboratory values, and mortality. Severity of disease was divided into mild (no oxygen therapy required), moderate (oxygen therapy administered through nasal canula or face mask), and severe (need of high-flow oxygen and/or ventilator). Laboratory values of interest were creatinine, estimated glomerular filtration rate (eGFR), potassium, C-reactive protein (CRP), blood lymphocytes, total leukocyte count, neutrophils, platelet count, cTnT, and NT-proBNP. Variables related to chloroquine treatment included date and length of treatment, and drug dosage (daily and total values).

For patients whose cardiac markers had not been analyzed during treatment or within 12 hours after their last dose of chloroquine, biobanked serum was sent for analysis of cTnT and NT-proBNP. Laboratory values from the control group and study group were matched as closely as possible with regard to time between onset of symptoms and collection of blood samples. Serum samples in the biobank that matched more closely in time were analyzed for cardiac markers. Definitions of heart failure (not likely, possible, or probable) were based on age-stratified reference levels from the Department of Clinical Chemistry at Sahlgrenska University Hospital with NT-proBNP < 300 (age 17 to 49 years), or < 400 (age > 49 years) indicating heart failure unlikely; NT-proBNP 300 to 400 (age 17 to 49 years), or 400 to 900 (age 50 to 75 years), or 400 to 1800 (age ≥ 76 years) indicating heart failure possible; and NT-proBNP > 400 (age 17 to 49 years), or > 900 (age 50 to 75 years), or > 1800 (age ≥ 76 years) indicating heart failure probable.

For patients who had an ECG performed during treatment or within two weeks of their last dose of chloroquine, a specialist in cardiology analyzed the ECGs focusing on arrhythmias and signs of heart dysfunction. When ECGs before and during or after treatment were available, they were analyzed and compared. For controls, ECGs taken as closely in time as possible to the cardiac markers were preferred; ECGs taken more than four weeks after onset of symptoms were excluded.
2.3 Statistics

Data was analyzed with IBM SPSS Statistics version 26, Stata software (version 16.1, StataCorp, College Station, Texas, USA), and R (version 4.1.0, R Core Team, 2020). Figures were produced using the R package ggplot2. Continuous variables are presented as median (I–III quartile), and categorical variables as absolute numbers (percentages). For comparisons between the two groups, the Mann Whitney U-test was used for continuous variables and the Fischer’s exact test for categorical variables. 

P-values less than 0.05 were considered statistically significant. We evaluated the relative importance of all variables from Tables 1–3, and the additional variables start of chloroquine treatment, admission to intensive care unit, day of blood test, activated partial thromboplastin time (APTT), international normalized ratio (INR), procalcitonin, magnesium, d-dimer, heart rate, heart rhythm, and normal QTc-time, for occurrence of significant cTnT leakage. Relative variable importance was calculated using a generalized boosted regression model as implemented in the R package gbm. Generalized boosted regression modelling is a highly efficient machine-learning method that handles a large number of observations and predictors and produces a prediction model in the form of an ensemble of weak prediction models. Typically, the decision trees allow for complex modeling of interactions and non-linear functions between predictors.

The study was approved by the Swedish Ethics Review Authority (Registration number 2020–01771). Patients were included after written informed consent to use data from their medical records and samples in research was obtained at the time of admission to the hospital.

| Table 1. Demographic characteristics of 60 study participants. | Total (n = 60) | Chloroquine (n = 20) | Controls (n = 40) |
|---------------------------------------------------------------|--------------|---------------------|------------------|
| Age; median (IQR)                                            | 60 (50–72)   | 61 (51–72)          | 60 (50–70)       |
| Gender; no (%)                                               |              |                     |                  |
| Male                                                         | 49 (82)      | 17 (85)             | 32 (80)          |
| Female                                                       | 11 (18)      | 3 (15)              | 8 (20)           |
| Body Mass Index; median (IQR)                                | 29 (26–32)   | 29 (27–31)          | 29 (26–32)       |
| Comorbidities, number; median (IQR)                          | 1 (0–3)      | 0 (0–3)             | 1 (1–3)          |
| Previous heart disease; no (%)                               | 12 (20)      | 5 (25)              | 7 (18)           |
| Ischemic heart disease; no (%)                               | 6 (10)       | 1 (5)               | 5 (13)           |
| Cardiovascular risk factors; no (%)                          | 34 (57)      | 9 (45)              | 25 (63)          |
| Other drugs affecting QTc time; no (%)                       | 5 (8)        | 2 (10)              | 3 (8)            |
| Severity of disease; no (%)                                  |              |                     |                  |
| Mild                                                         | 3 (5)        | 1 (5)               | 2 (5)            |
| Moderate                                                     | 12 (20)      | 4 (20)              | 8 (20)           |
| Severe                                                       | 45 (75)      | 15 (75)             | 30 (75)          |
| Ventilator; no (%)                                           | 36 (60)      | 13 (65)             | 23 (58)          |
| Deceased; no (%)                                             | 11 (18)      | 3 (15)              | 8 (20)           |
| Chloroquine treatment length, days; median (IQR)             | 5 (3–8)      | 5 (3–8)             | 0                |
| Total dosage chloroquine, mg (IQR)                           | 5250 (3000–7500) | 5250 (3000–7500) | 0                |

IQR = interquartile range.

* Data available for 42 patients.

 Data available for 14 patients.

 Data available 28 patients.

 Cardiovascular risk factors: hypertension, diabetes, dyslipidemia.

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All data were retrospectively collected between April and August 2020. The data was not anonymized when compiled by the authors, but was anonymized before statistical analysis.

3. Results

In the six weeks between 26 February and 15 April 2020, a total of 226 patients were admitted to the Department of Infectious Diseases, Sahlgrenska University Hospital with a diagnosis of COVID-19. Twenty-five of them were administered chloroquine, five of whom were excluded from our study because their cardiac markers had not been analyzed or no stored serum samples were available from their treatment period. The remaining 20 patients comprised the chloroquine arm. Forty matched controls who had not undergone chloroquine treatment were also included.

The groups were demographically similar, with a median (interquartile range [IQR]) age of 61 (51–72) years in the study group, compared with a median age (IQR) of 60 (50–70) years in the control group (Table 1). In the study group there were 17 (85%) men, while there were 32 (80%) men in the control group. The median body mass index (BMI) was 29 in both groups.

Five (25%) patients in the study group and eight (18%) in the control group had a history of...
heart disease. Ischemic heart disease was present in one (14%) patient with elevated cTnT in the study group, whereas four patients (57%) had a cardiovascular risk factor (diabetes, hypertension and/or hyperlipidemia). In the control group there were four (17%) patients with elevated cTnT and ischemic heart disease, and 16 (70%) patients with a cardiovascular risk factor. Two (10%) patients in the study group and three (8%) in the control group were taking medication known to potentially affect QT-interval (2 alfuzosin, 2 escitalopram, 1 tacrolimus). There were similar rates of disease severity in both groups: 5% with mild, 20% with moderate, and 75% with severe COVID-19. Among patients receiving chloroquine, 13 (65%) required invasive mechanical ventilation compared to 23 (58%) of the controls. The overall in-hospital mortality rate was 18% \((n = 11)\) with three deaths occurring in the study group and eight in the control group.

Median (IQR) duration of treatment with chloroquine was 5 (3–8) days, and the median total dosage was 5250 (3000–7500) mg. Time between onset of symptoms and first day of treatment was a median (IQR) of 10 (8–13) days.

The details of the biochemical analyses are presented in Table 2. Median (IQR) time between symptom onset and sampling of cardiac biomarkers was 14 (10–15) days in the study group and 12 (10–14) days in the control group. All other biochemical variables were recorded as closely as possible in time to that of the cardiac markers. There were no statistically significant differences in any of the variables analyzed. For patients receiving chloroquine, cTnT levels were in median (IQR) 10.8 (7.4–22.8) ng/L, and 17.9 (8.0–55.1) ng/L for the controls. Seven (35%) patients in the study group and 23 (58%) controls had elevated cTnT \(> 13.9\) ng/L. Fig 1 shows the relative importance of several variables in predicting elevation in cTnT. Treatment with chloroquine had a low relative importance, whereas eGFR, neutrophils, comorbidity, and APPT had high relative importance on cTnT levels.

Median (IQR) NT-proBNP was 399 (143–1235) ng/L in the study group and 349 (130–765) ng/L in the control group. Ten (50%) patients had NT-proBNP values that did not indicate heart failure, and 10 (50%) patients in the study group had possible or probable heart failure. There were 23 (58%) patients with no heart failure, and 17 (43%) patients with possible or probable heart failure, in the control group.

ECGs were performed in ten patients on chloroquine treatment and in 31 controls (Table 3). There were no significant differences in QTc interval prolongation, ST-segment abnormalities, or T-wave inversion. Three patients had their ECGs performed before and after initiation of chloroquine. In one of these patients, QTc interval went from normal to pathologic, increasing from 456 milliseconds (ms) to 473 ms. In the other two, the QTc interval remained the same in one and increased by five ms in the other patient, none of the increases being pathologic. Abnormalities in ST-segment were not present in any of the ECGs. However, T-wave inversion appeared in two patients after initiation of treatment.

4. Discussion

We have examined biomarkers of ischemic heart injury and heart failure in patients with varying severities of COVID-19 who either received or did not receive treatment with chloroquine phosphate. In this small study population, we found no significant differences in cTnT or NT-proBNP levels between patients who received chloroquine phosphate and those who did not, indicating that chloroquine had no general effect on cardiac biomarkers.

The COVID-19 outbreak is a global pandemic with considerable mortality. Effective and well-tolerated treatments are therefore urgently needed. The old anti-malarial drug chloroquine phosphate was one of the first pharmaceuticals to be widely used against COVID-19, based on positive preliminary reports from China and France [12, 15]. It was immediately
adopted in most countries, including Sweden. However, doubts were soon raised about the safety of the drug. The Department of Infectious Diseases at Sahlgrenska University Hospital was among the first clinics to stop using chloroquine at the end of March 2020, because of the lack of evidence that it was effective and its potential for causing severe adverse events. In early April, a small French study [21] could not demonstrate the positive effects on viral clearance that had been reported earlier [15]. Doubts about using chloroquine to treat COVID-19 patients were soon raised because of reports of severe adverse events and inconclusive data on clinical effects [22]. Descriptions of the adverse effects of chloroquine became more frequent and several countries discontinued its use as a standard treatment for COVID-19. There were at least 11 active (recruiting or not-yet recruiting) clinical trials of chloroquine as a treatment for COVID-19 (prophylactic or treatment of active disease) registered at clinicaltrials.gov on 18 June 2021 [23]. However, several clinical trials were halted as the positive preliminary reports of its efficacy failed to be confirmed in several studies [18, 20]. Chloroquine treatment of COVID-19 is no longer supported by most clinicians. Nevertheless, it may well be of interest in future pandemics/virus outbreaks given that it has had historical interest for other infectious diseases such as SARS [24], influenza [25], yellow fever [26], and HIV [27].

Fig 1. Relative importance of predictors of cardiovascular involvement. eGFR = estimated glomerular filtration rate, comorbidity = numbers of comorbidities, APTT = activated partial thromboplastin time, INR = international normalized ratio, T_wave_invers = T-wave inversion, Chlor_phosph_days = number of days with chloroquine phosphate, CV_risk_factor = cardiovascular risk factors (hypertension, diabetes, dyslipidemia), Chlor_phosph_init = days since symptom onset when chloroquine phosphate was initiated, Blood_test_day = day since symptom onset when blood samples were collected, BMI = body mass index, Chlorophosph = treatment with chloroquine phosphate, Heart_disease = type of previous heart disease specified (cardiac arrhythmia 7, valvular heart disease 6, ischemic heart disease 6, congestive heart failure 1), Chlor_phosph_dose = total dose of chloroquine phosphate. Missing data: eGFR 1, APTT 21, thrombocytes 6, heart rate 19, INR 21, potassium 1, T wave inversion 19, CRP 2, BMI 18, magnesium 34, procalcitonin 34, QTc time 19, QT time 19, d-dimer 33, ST abnormalities 19, rhythm 19. Relative variable importance was calculated using a generalized boosted regression model.

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To the best of our knowledge, ours is the first study to analyze markers of heart failure and ischemic injury in patients treated with chloroquine phosphate for COVID-19. A pre-pandemic review from 2018 reported that 8/127 patients receiving long-term treatment with hydroxychloroquine and/or chloroquine had elevated cTnT, and 21/46 patients had an ejection fraction less than 40%, indicating mild-to-severe heart failure [28]. In our study, we could find no increased risk of elevated cTnT among patients who received chloroquine phosphate as compared to those who did not, and the slightly elevated NT-proBNP levels indicating possible or probable heart failure (50% in the study group versus 43% in the control group) were not statistically significant. Moreover, our analyses of ECGs showed no significant differences in signs of ischemic heart injury. Treatment duration in the pre-pandemic study was approximately seven to eight years longer than in our study, and the risk of heart failure may increase with long-term treatment. Thus far, there is no evidence of increased risk for ischemic heart injury during short-term treatment with chloroquine phosphate, and the risk for ischemic injury over long-term treatment appears to be only slightly elevated.

Several studies have shown an increased risk of cardiovascular events during lower respiratory tract infections [29, 30], and specifically during infection with SARS-CoV-2 [3, 31], a finding supported by the high number of patients with elevated cTnT and NT-proBNP in both groups in our study. In a machine learning-based analysis of predictors of elevated cTnT, we found that chloroquine had a low relative importance, whereas factors such as kidney function, neutrophils, comorbidity, and APTT had high relative importance. Age and thrombocytes were of moderate relative importance. Elevated creatinine at admission was found to be a predictor of myocardial injury in COVID-19 in another retrospective analysis [32], and studies have shown that patients with chronic kidney disease generally have higher cTnT levels [33]. Increased APTT, coagulopathy, inflammation, and advanced age have previously been shown to be associated with myocardial injury during COVID-19 infection [34, 35]. Risk factors for cardiovascular disease were present in most patients with elevated cTnT levels, a finding supported by a previous study in which myocardial injury was seen to be significantly more common in COVID-19 patients with chronic coronary syndromes than in patients without [36]. Therefore, while it is difficult to distinguish the cause of myocardial injury among our participants, it is recommendable that chloroquine be used with caution specifically in patients with previous cardiovascular disease, impaired kidney function, and advanced age, until further research has been done.

The pharmacological effects of chloroquine phosphate in cardiac toxicity that have been recognized thus far involve an impact on the electrophysical properties of the heart, such as blocking potassium channels, which in turn could cause drug-induced long QT-syndrome [9]. In the pre-pandemic review cited earlier [28], electrophysical conduction disorders were reported in 85% of patients receiving hydroxychloroquine and/or chloroquine for various inflammatory and infectious diseases. Of 91 patients with COVID-19 receiving hydroxychloroquine and azithromycin, 23% had significant QTc prolongation in a retrospective study [37]. This has been confirmed in further studies, that included individuals with suspected torsade de points [16]. Researchers studying patients with COVID-19 receiving hydroxychloroquine as monotherapy found that 19% developed prolonged QTc of 500 ms and 3% had an increase in QTc—time of 60 ms [17]. A retrospective study of 397 patients with COVID-19 treated with chloroquine [38] showed that the QTc interval gradually increased during the treatment period, probably due to the drug’s long half-life of 1.6 days. On the contrary, a multicenter study of 649 COVID-19 patients receiving hydroxychloroquine found only a modest QTc prolongation [39]. Factors identified as predictors of prolonged QTc-time during treatment with the combination hydroxychloroquine and azithromycin include increased troponin, older age, congestive heart failure, and higher admission creatinine [40]. In our study, where ECGs
unfortunately were only available for 41 patients, we could not see any significant differences in QTc prolongation or signs of ischemic heart injury when comparing patients with and without chloroquine treatment. However, one of the three patients with a baseline ECG, and a second ECG taken while treated for COVID-19 developed a pathologic QTc prolongation, indicating a potential risk of severe arrhythmia in using chloroquine to treat COVID-19.

Our study has several limitations. One major limitation is the small sample size, which makes statistical comparisons uncertain. As data was collected retrospectively, we were unable to measure serum concentrations of chloroquine phosphate. Since there was no systematic ECG collection, we did not have baseline and follow-up ECGs for most of the patients studied.

5. Conclusions

We found no differences in acute ischemic heart injury or heart failure in our relatively small sample of hospitalized patients with COVID-19, whether they received treatment with chloroquine phosphate or not. ECG findings from the even smaller number of patients who had ECGs performed showed no differences regarding QTc prolongation or signs of ischemic heart injury. However, in the absence of large-scale studies, chloroquine treatment for COVID-19 infection cannot be considered safe with regard to myocardial infarction and heart failure. Therefore, it cannot be recommended at the present time that chloroquine be used to treat COVID-19 outside of clinical trials, considering the lack of evidence of its efficacy and the elevated risk it presents for arrhythmia and sudden cardiac death.

Author Contributions

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References

1. Wu Z, McGoogan JM. Characteristics of and Important Lessons From the Coronavirus Disease 2019 (COVID-19) Outbreak in China. Summary of a Report of 72314 Cases From the Chinese Center for Disease Control and Prevention. JAMA. 2020; 323(13):1239–42. https://doi.org/10.1001/jama.2020.2648 PMID: 32091533

2. Oran DP, Topol EJ. Prevalence of Asymptomatic SARS-CoV-2 Infection: A Narrative Review. Ann Intern Med. 2020; 173(5):362–7. https://doi.org/10.7326/M20-3012 PMID: 32491919

3. Wang D, Hu B, Hu C, Zhu F, Liu X, Zhang J, et al. Clinical Characteristics of 138 Hospitalized Patients With 2019 Novel Coronavirus-Infected Pneumonia in Wuhan, China. JAMA. 2020; 323(11):1061–9. https://doi.org/10.1001/jama.2020.1585 PMID: 32031570

4. Phua J, Weng L, Ling L, Egi M, Lim CM, Divatia JV, et al. Intensive care management of coronavirus disease 2019 (COVID-19): challenges and recommendations. Lancet Respir Med. 2020; 8(5):506–17. https://doi.org/10.1016/S2213-2600(20)30161-2 PMID: 32272080

5. Intensivvårdsregistret S. Svenska Intensivvårdsregistret [cited 2020 December 1]. https://www.icuregswe.org/data—resultat/covid-19-i-svensk-intensivvard/.
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6. EMA. Treatments and vaccines for COVID-19 [cited 2021 January 24]. https://www.ema.europa.eu/en/human-regulatory/overview/public-health-threats/coronavirus-disease-covid-19/treatments-vaccines-covid-19.

7. FDA. Coronavirus disease 2019 (COVID-19), Resources for Health Professionals; FDA; [cited 2021 January 24]. https://www.fda.gov/health-professionals/coronavirus-disease-2019-covid-19-resources-health-professionals.

8. Jean S-S, Hsueh P-R. Old and re-purposed drugs for the treatment of COVID-19. Expert Rev Anti Infect Ther. 2020;1–5. https://doi.org/10.1080/14787210.2020.1771181 PMID: 32419524

9. Kamp TJ, Hamdan MH, January CT. Chloroquine or Hydroxychloroquine for COVID-19: Is Cardiotoxicity a Concern? J Am Heart Assoc. (2047–9980 (Electronic)). https://doi.org/10.1161/JAHA.120.016887 PMID: 32463308

10. Schrezenmeier E, Dorner T. Mechanisms of action of hydroxychloroquine and chloroquine: implications for rheumatology. Nat Rev Rheumatol. 2020; 16(3):155–66. https://doi.org/10.1038/s41584-020-0372-x PMID: 32034323

11. Rainsford KD, Parke AL, Clifford-Rashotte M, Kean WF. Therapy and pharmacological properties of hydroxychloroquine and chloroquine in treatment of systemic lupus erythematosus, rheumatoid arthritis and related diseases. Inflammopharmacology. 2015; 23(5):231–69. https://doi.org/10.1007/s10787-015-0239-y PMID: 26246385

12. Zhou D, Dai SM, Tong Q. COVID-19: a recommendation to examine the effect of hydroxychloroquine in patients with COVID-19. BioMed Research International. 2021; 2021:1912497. https://doi.org/10.1155/2021/1912497

13. Liu J, Cao R, Xu M, Wang X, Zhang H, Hu H, et al. Hydroxychloroquine, a less toxic derivative of chloroquine, is effective in inhibiting SARS-CoV-2 infection in vitro. Cell Discov. 2020; 6:16.

14. Gao J, Tian Z, Yang X. Breakthrough: Chloroquine phosphate has shown apparent efficacy in treatment of COVID-19 associated pneumonia in clinical studies. Biosci Trends. 2020; 14(1):72–3. https://doi.org/10.5582/bst.2020.101663 PMID: 32289548

15. Gautret P, Lagier JC, Parola P, Hoang VT, Meddeb L, Sevestre J, et al. Chloroquine and hydroxychloroquine in the treatment of patients hospitalised with COVID-19: a multicentre, observational study. Lancet Infect Dis. 2020; 20(5):513–9. https://doi.org/10.1016/S1473-3099(20)30086-9

16. Cipriani A, Zorzi A, Ceccato D, Capone F, Parolin M, Donato F, et al. Arrhythmic profile and 24-hour QT interval variability in COVID-19 patients treated with Hydroxychloroquine and azithromycin. Int J Cardiol. 2020; 316:280–4. https://doi.org/10.1016/j.ijcard.2020.05.036 PMID: 32439366

17. Mercuro NJ, Yen CF, Shim DJ, Maher TR, McCoy CM, Zimetbaum PJ, et al. Risk of QT Interval Prolongation Associated With Use of Hydroxychloroquine With or Without Concomitant Azithromycin Among Hospitalized Patients Testing Positive for Coronavirus Disease 2019 (COVID-19). JAMA Cardiol. 2020; 5(9):1036–41. https://doi.org/10.1001/jamacardio.2020.1834 PMID: 32936252

18. Mitja O, Corbacho-Monne M, Ubals M, Tebe C, Penafiel J, Tobias A, et al. Hydroxychloroquine for Early Treatment of Adults with Mild Covid-19: A Randomized-Controlled Trial. Clin Infect Dis. 2020;Epub ahead of print. https://doi.org/10.1093/cid/ciaa1009 PMID: 32674126

19. WHO. “Solidarity” clinical trial for COVID-19 treatments 2020 [https://www.who.int/emergencies/diseases/novel-coronavirus-2019/global-research-on-novel-coronavirus-2019-ncov/solidarity-clinical-trial-for-covid-19-treatments].

20. Self WH, Semler MW, Leither LM, Casey JD, Angus DC, Brower RG, et al. Effect of Hydroxychloroquine on Clinical Status at 14 Days in Hospitalized Patients With COVID-19: A Randomized Clinical Trial. JAMA. 2020; 324(21):2165–76. https://doi.org/10.1001/jama.2020.22240 PMID: 33156621

21. Molina JM, Delaugerre C, Le Goff J, Mela-Lima B, Ponscarne D, Goldwirt L, et al. No evidence of rapid antiviral clearance or clinical benefit with the combination of hydroxychloroquine and azithromycin in patients with severe COVID-19 infection. Med Mal Infect. 2020; 50(4):384. https://doi.org/10.1016/j.medmal.2020.03.006 PMID: 32240719

22. Yazdany J, Kim AHJ. Use of Hydroxychloroquine and Chloroquine During the COVID-19 Pandemic: What Every Clinician Should Know. Ann Intern Med. 2020; 172(11):754–5. https://doi.org/10.7326/M20-1334 PMID: 32232419

23. ClinicalTrials.gov: U.S. National Library of Medicine; [cited 2021 18 June]. https://clinicaltrials.gov/ct2/results?cond=Covid19&term=chloroquine&cntry=&state=&city=&dist*&HYPERLINK*https://clinicaltrials.gov/ct2/results?cond=Covid19&term=chloroquine&cntry=&state=&city=&dist*&HYPERLINK*https://clinicaltrials.gov/ct2/results?cond=Covid19&term=chloroquine&cntry=&state=&city=&dist*&HYPERLINK
Cardiac biomarkers in patients with COVID-19 receiving chloroquine phosphate

24. Vincent MJ, Bergeron E, Benjannet S, Erickson BR, Rollin PE, Ksiazek TG, et al. Chloroquine is a potent inhibitor of SARS coronavirus infection and spread. Virol J 2005; 2:69. https://doi.org/10.1186/1743-422X-2-69 PMID: 16115318

25. Paton NI, Lee L, Xu Y, Ooi EE, Cheung YB, Archuleta S, et al. Chloroquine for influenza prevention: a randomised, double-blind, placebo controlled trial. Lancet Infect Dis. 2011; 11(9):677–83. https://doi.org/10.1016/S1473-3099(11)70065-2 PMID: 21550310

26. Miller BR, Tsai TF, Mitchell CJ. Aedes aegypti and yellow fever virus: the effect of chloroquine on infection and transmission rates. Trans R Soc Trop Med Hyg. 1987; 81(1):111–2. https://doi.org/10.1016/0035-9020(87)90298-7 PMID: 3445296

27. Sperber K, Louie M, Kraus T, Proner J, Sapira E, Lin S, et al. Hydroxychloroquine treatment of patients with human immunodeficiency virus type 1. Clin Ther. 1995; 17(4):622–36. https://doi.org/10.1016/0149-2918(95)80039-5 PMID: 8565026

28. Chatre C, Roubille F, Vernhet H, Jorgensen C, Pers YM. Cardiac Complications Attributed to Chloroquine and Hydroxychloroquine: A Systematic Review of the Literature. Drug Saf. 2018; 41(10):919–31. https://doi.org/10.1007/s40262-018-0689-4 PMID: 29858838

29. Smeeth L, Thomas SI Fau—Hall AJ, Hall Aj Fau—Hubbard R, Hubbard R Fau—Farrington P, Farrington P. Fau—Vallance P, Vallance P. Risk of myocardial infarction and stroke after acute infection or vaccination. N Engl J Med. (1533–4406 (Electronic)). https://doi.org/10.1111/CIRCULATIONAHA.111.040766 PMID: 2219349

30. Zhou F, Yu T, Du R, Fan G, Liu Y, Li Z, et al. Clinical course and risk factors for mortality of adult inpatients with COVID-19 in Wuhan, China: a retrospective cohort study. Lancet (London, England). 2020; 395(10229):1054–62. https://doi.org/10.1016/S0140-6736(20)30566-3 PMID: 32171076

31. Efros O, Barda N, Meisel E, Leibowitz A, Fardman A, Rahav G, et al. Myocardial injury in hospitalized patients with COVID-19 infection: Risk factors and outcomes. PLoS One. 2021; 16(2):e0247800. https://doi.org/10.1371/journal.pone.0247800 PMID: 33635914

32. Kraus D, von Jeinsen B, Tzikas S, Palapies L, Zeller T, Bickel C, et al. Cardiac Troponins for the Diagnosis of Acute Myocardial Infarction in Chronic Kidney Disease. J Am Heart Assoc. 2018; 7(19):e008032. https://doi.org/10.1161/JAHA.111.008032 PMID: 30371308

33. Fan ZX, Yang J, Zhang J, He C, Wu H, Yang CJ, et al. Analysis of influencing factors related to elevated serum troponin I level for COVID-19 patients in Yichang, China. Cardiovasc Diagn Ther. 2020; 10(4):678–86. https://doi.org/10.21037/cdt-20-510 PMID: 32968624

34. Arevalos V, Ortega-Paz L, Rodriguez-Arias JJ, Calvo M, Castrillo L, Salazar A, et al. Myocardial Injury in COVID-19 Patients: Association with Inflammation, Coagulopathy and In-Hospital Prognosis. J Clin Med. 2021; 10(10). https://doi.org/10.3390/jcm10103263 PMID: 33053826

35. Marag I, Hummel JA-O, Tautel R, Chamoun R, Workman V, Li C, et al. Incidence and Determinants of QT Interval Prolongation in COVID-19 Patients Treated with Hydroxychloroquine and Azithromycin. J Cardiovasc Electrophysiol. (1540–8167 (Electronic)). https://doi.org/10.1111/jce.14594 PMID: 32485061

36. Sinkeler FS, Berger FA, Muntinga HJ, Jansen M. The risk of QTc-interval prolongation in COVID-19 patients treated with chloroquine. Neth Heart J. 28(7–8):418–23. https://doi.org/10.1007/s12471-020-01462-6 PMID: 32648153

37. Gasperetti A, Biffi M, Dufu F, Schiavone M, Ziacchi M, Mitacchione G, et al. Arrhythmic safety of hydroxychloroquine in COVID-19 patients from different clinical settings. Europace. 2020; 22(12):1855–63. https://doi.org/10.1093/europace/euaa216 PMID: 32971536

38. O’Connell TF, Bradley CJ, Abbas AE, Williamson BD, Rusia A, Tawney AM, et al. Hydroxychloroquine/Azithromycin Therapy and QT Prolongation in Hospitalized Patients With COVID-19. JACC Clin Electrophysiol. 2021; 7(1):16–25. https://doi.org/10.1016/j.jacep.2020.07.016 PMID: 33478708