

Mortality outcomes with hydroxychloroquine and chloroquine in COVID-19: an international collaborative meta-analysis of randomized trials

Cathrine Axfors MD, PhD^{a,1,2}, Andreas M. Schmitt MD^{a,3,4}, Perrine Janiaud PhD³, Janneke van 't Hooft MD, PhD^{1,5}, Sherief Abd-Elsalam MD, PhD⁶, Ehab F. Abdo MD, PhD⁷, Benjamin S. Abella MD, MPhil⁸, Javed Akram MBBS, FRCP⁹, Ravi K. Amaravadi MD¹⁰, Derek C. Angus MD^{11,12}, Yaseen M. Arabi MD, FCCP, FCCM¹³, Shehnoor Azhar BDS, MPH¹⁴, Lindsey R. Baden MD¹⁵, Arthur W. Baker MD, MPH¹⁶, Leila Belkhir MD, PhD¹⁷, Thomas Benfield MD, DMSc¹⁸, Marvin A.H. Berrevoets MD¹⁹, Cheng-Pin Chen MD²⁰, Tsung-Chia Chen MD²¹, Shu-Hsing Cheng MD, PhD²⁰, Chien-Yu Cheng MD²⁰, Wei-Sheng Chung MD, PhD²¹, Yehuda Z. Cohen MD²², Lisa N. Cowan MS²², Olav Dalgard MD, PhD^{23,24}, Fernando F. de Almeida e Val PhD²⁵, Marcus V.G. de Lacerda PhD^{25,26}, Gisely C. de Melo PhD^{25,27}, Lennie Derde MD, PhD^{28,29}, Vincent Dubee MD, PhD³⁰, Anissa Elfakir MSc³¹, Anthony C. Gordon MD³², Carmen M. Hernandez-Cardenas MD, MSc³³, Thomas Hills DPhil^{34,35}, Andy I.M. Hoepelman Prof. Dr.³⁶, Yi-Wen Huang MD³⁷, Bruno Igau PhD²², Ronghua Jin MD, PhD³⁸, Felipe Jurado-Camacho MD, MSc³³, Khalid S. Khan MBBS, MRCOG³⁹, Peter G Kremsner MD^{40,41,42}, Benno Kreuels MD^{43,44}, Cheng-Yu Kuo MD⁴⁵, Thuy Le MD, PhD¹⁶, Yi-Chun Lin MD²⁰, Wu-Pu Lin MD⁴⁶, Tse-Hung Lin MD³⁷, Magnus Nakrem Lyngbakken MD, PhD^{47,24}, Colin McArthur MBChB^{34,35,48}, Bryan J. McVerry MD⁴⁹, Patricia Meza-Meneses MD⁵⁰, Wuelton M. Monteiro PhD^{25,27}, Susan C. Morpeth PhD⁵¹, Ahmad Mourad MD⁵², Mark J. Mulligan MD, FIDSA^{53,54}, Srinivas Murthy MD, MHSc⁵⁵, Susanna Naggie MD¹⁶, Shanti Narayanasamy MBBS¹⁶, Alistair Nichol PhD^{48,56,57,58}, Lewis A. Novack MS⁵⁹, Sean M. O'Brien PhD⁶⁰, Nwora Lance Okeke MD, MPH¹⁶, Léna Perez MS⁶¹, Rogelio Perez-Padilla MD⁶², Laurent Perrin MSc⁶³, Arantxa Remigio-Luna MD⁶², Norma E. Rivera-Martinez MD, MSc⁶⁴, Frank W. Rockhold PhD⁶⁰, Sebastian Rodriguez-Llamazares MD, MPH⁶², Robert Rolfe MD¹⁶, Rossana Rosa MD⁶⁵, Helge Røsjø MD, PhD^{66,24}, Vanderson S. Sampaio PhD^{25,67}, Todd B. Seto MD, MPH^{68,69}, Muhammad Shehzad PhD⁷⁰, Shaimaa Soliman MD, PhD⁷¹, Jason E. Stout MD, MHS⁷⁶, Ileri Thurion-Romero MD, MSc⁶², Andrea

B. Troxel ScD⁷², Ting-Yu Tseng MD²¹, Nicholas A. Turner MD, MHSc¹⁶, Robert J. Ulrich MD⁷³, Stephen R. Walsh MD¹⁵, Steve A. Webb MD, MPH, PhD^{48,74}, Jesper M. Weehuizen MD³⁶, Maria Velinova MD, PhD⁷⁵, Hon-Lai Wong MD⁷⁶, Rebekah Wrenn PharmD¹⁶, Fernando G. Zampieri MD, PhD^{77,78,79}, Wu Zhong PhD⁸⁰, David Moher PhD⁸¹, Steven N. Goodman MD, PhD^{1,82,83}, John P.A. Ioannidis MD, DSc^{1,82,83,84,85}, Lars G. Hemkens MD, MPH*^{1,3,85}

^aEqual contributions, ¹Meta-Research Innovation Center at Stanford (METRICS), Stanford University, Stanford, CA, USA, ²Department for Women's and Children's Health, Uppsala University, Uppsala, Sweden, ³Department of Clinical Research, University Hospital Basel, University of Basel, Basel, Switzerland, ⁴Department of Medical Oncology, University of Basel, Basel, Switzerland, ⁵Amsterdam University Medical Center, Amsterdam University, Amsterdam, the Netherlands, ⁶Tropical Medicine and Infectious Diseases Department, Faculty of Medicine, Tanta University, Tanta, Egypt, ⁷Tropical Medicine and Gastroenterology Department, Faculty of Medicine, Assiut University, Assiut, Egypt, ⁸Department of Emergency Medicine, University of Pennsylvania, USA, ⁹Department of Internal Medicine, Vice Chancellor, University of Health Sciences, Khayaban e Jamia Punjab, Lahore, Punjab, 54000, Pakistan, ¹⁰Abramson Cancer Center and Department of Medicine, University of Pennsylvania, Philadelphia, USA, ¹¹Department of Critical Care Medicine, The Clinical Research Investigation and Systems Modeling of Acute Illness (CRISMA) Center, University of Pittsburgh, Pittsburgh, Pennsylvania, USA, ¹²the UPMC Health System Office of Healthcare Innovation, University of Pittsburgh Medical Centre, Pittsburgh, Pennsylvania, USA, ¹³Intensive Care Department, King Saud Bin Abdulaziz University for Health Sciences and King Abdullah International Medical Research Center, Riyadh, Saudi Arabia, ¹⁴Department of Public Health, University of Health Sciences, Khayaban e Jamia Punjab, Lahore, Punjab, 54000, Pakistan, ¹⁵Division of Infectious Diseases, Brigham and Women's Hospital, Boston, MA, USA, ¹⁶Department of Medicine, Division of Infectious Diseases and International Health, Duke University Medical Center, Durham, NC, 27710, USA, ¹⁷Infectious Diseases Department, Cliniques universitaires Saint-Luc, Université Catholique de Louvain, Brussels, Belgium, ¹⁸Center of Research & Disruption of Infectious Diseases, Department of Infectious Diseases, Copenhagen University Hospital, Amager

and Hvidovre, Kettegaard Alle 30, Hvidovre, 2650, Denmark, ¹⁹Department of Internal Medicine, Elisabeth-Tweesteden hospital, Tilburg, the Netherlands, ²⁰Department of Infectious Diseases, Taoyuan General Hospital, Ministry of Health and Welfare, Taoyuan, 330, Taiwan, ²¹Department of Internal Medicine, Taichung Hospital, Ministry of Health and Welfare, Taichung, Taiwan, ²²Sanofi, Bridgewater, NJ, USA, ²³Department of Infectious Diseases, Division of Medicine, Akershus University Hospital, Lørenskog, Norway, ²⁴Institute of Clinical Medicine, Faculty of Medicine, University of Oslo, Oslo, Norway, ²⁵Fundação de Medicina Tropical Dr. Heitor Vieira Dourado, Pedro Teixeira, 25, Manaus, AM, 69040-000, Brazil, ²⁶Instituto Leonidas e Maria Deane – ILM, FIOCRUZ-AM, Teresina, 476, Manaus, AM, 69.057-070, Brazil, ²⁷Universidade do Estado do Amazonas, Djalma Batista, 2470, Manaus, AM, 69050-300, Brazil, ²⁸Julius Center for Health Science and Primary Care, University Medical Center Utrecht, Utrecht, Netherlands, ²⁹Intensive Care Centre, University Medical Center Utrecht, Utrecht, Netherlands, ³⁰Infectious and Tropical Diseases Department, Angers University Hospital, Angers, France, ³¹Ividata Life Sciences, Levallois-Perret, France, ³²Department of Surgery and Cancer, Anaesthetics, Pain Medicine, and Intensive Care Medicine, Imperial College London and Imperial College Healthcare NHS Trust, London, United Kingdom, ³³Critical Care Department, Instituto Nacional de Enfermedades Respiratorias Ismael Cosío Villegas, Mexico, ³⁴Medical Research Institute of New Zealand, Wellington, New Zealand, ³⁵Auckland City Hospital, Auckland, New Zealand, ³⁶Department of Infectious Diseases, University Medical Center Utrecht, Utrecht, the Netherlands, ³⁷Department of Internal Medicine, Chang Hua Hospital, Ministry of Health and Welfare, Changhua, Taiwan, ³⁸Beijing Youan Hospital, Capital Medical University, Beijing, P. R. China, ³⁹Department of Preventive Medicine & Public Health, University of Granada, Hospital Real, Avenida del Hospicio, Granada, Granada, 18010, Spain, ⁴⁰Institute of Tropical Medicine, University of Tübingen, Tübingen, Germany, ⁴¹Centre de Recherches Médicales de Lambaréné, Lambaréné, Gabon, ⁴²German Center for Infection Research, Partner Site Tübingen, Tübingen, Germany, ⁴³Department of Medicine, Division of Tropical Medicine and Division of Infectious Diseases, University Medical Center Hamburg-Eppendorf, Hamburg, Germany, ⁴⁴Department of Tropical Medicine, Bernhard Nocht Institute for Tropical Medicine, Hamburg, Germany, ⁴⁵Department of Internal Medicine, Pingtung Hospital, Ministry of Health and Welfare,

Pingtung, Taiwan, ⁴⁶Department of Internal Medicine, Taipei Hospital, Ministry of Health and Welfare, New Taipei City, Taiwan, ⁴⁷Division of Medicine, Akershus University Hospital, Lørenskog, Norway, ⁴⁸School of Epidemiology and Preventive Medicine, Australian and New Zealand Intensive Care Research Centre, Monash University, Melbourne, Victoria, Australia, ⁴⁹Department of Medicine, University of Pittsburgh, Pittsburgh, Pennsylvania, USA, ⁵⁰Hospital Regional de Alta especialidad de Ixtapaluca, Mexico, ⁵¹Middlemore Hospital, Auckland, New Zealand, ⁵²Department of Medicine, Duke University Medical Center, Durham, NC, 27710, USA, ⁵³Department of Microbiology, NYU Grossman School of Medicine, 550 First Ave., New York, NY, 10016, USA, ⁵⁴Department of Internal Medicine, Division of Infectious Diseases and Immunology, NYU Grossman School of Medicine, 550 First Ave., New York, NY, 10016, USA, ⁵⁵University of British Columbia School of Medicine, Vancouver, British Columbia, Canada, ⁵⁶Department of Intensive Care, Alfred Health, Melbourne, Victoria, Australia, ⁵⁷Department of Anesthesia and Intensive Care, St Vincent's University Hospital, Dublin, Ireland, ⁵⁸School of Medicine and Medical Sciences, University College Dublin, Dublin, Ireland, ⁵⁹Division of Infectious Diseases, Brigham and Women's Hospital, Harvard Medical School, Boston, MA, USA, ⁶⁰Department of Biostatistics and Bioinformatics, Duke University Medical Center and Duke Clinical Research Institute, Durham, NC, 27710, USA, ⁶¹Excelya, Montpellier, France, ⁶²Department of Smoking and COPD, Instituto Nacional de Enfermedades Respiratorias Ismael Cosío Villegas, Mexico, ⁶³Sanofi, Montpellier, France, ⁶⁴Hospital Regional de Alta especialidad de Oaxaca, Mexico, ⁶⁵UnityPoint Health, Des Moines, IA, 50309, USA, ⁶⁶Division of Research and Innovation, Akershus University Hospital, Lørenskog, Norway, ⁶⁷Fundação de Vigilância em Saúde do Amazonas, Djalma Batista, 2470, Manaus, AM, 69093-018, Brazil, ⁶⁸University of Hawaii John A. Burns School of Medicine, HI, USA, HI, USA, ⁶⁹The Queen's Medical Center, Honolulu, HI, USA, ⁷⁰Department of Pharmacology, University of Health Sciences, Khayaban e Jamia Punjab, Lahore, Punjab, 54000, Pakistan, ⁷¹Public health and Community Medicine, Menoufia University, Menoufia, Egypt, ⁷²Department of Population Health, Division of Biostatistics, NYU Grossman School of Medicine, 180 Madison Avenue, Fifth Floor, 5-55, New York, NY, 10016, USA, ⁷³Department of Medicine, Division of Infectious Diseases and Immunology, NYU Grossman School of Medicine, 550 First Ave., New York, NY, 10016, USA, ⁷⁴St. John of God Hospital,

Subiaco, Western Australia, Australia, ⁷⁵PRA Health Science, Groningen, Netherlands, ⁷⁶Department of Internal Medicine, Keelung Hospital, Ministry of Health and Welfare, Keelung, Taiwan, ⁷⁷Research Institute, HCor-Hospital do Coração, São Paulo, São Paulo, Brazil, ⁷⁸Research Institute, BRICNet - Brazilian Research in Intensive Care Network, São Paulo, São Paulo, Brazil, ⁷⁹IDor Research Institute, São Paulo, São Paulo, Brazil, ⁸⁰National Engineering Research Center for the Emergency Drug, Beijing Institute of Pharmacology and Toxicology, Beijing, P. R. China, ⁸¹Centre for Journalology, Clinical Epidemiology Program, Ottawa Hospital Research Institute, 501 Smyth Road, Ottawa K1H 8L6, Canada, ⁸²Stanford University School of Medicine, Stanford, CA, USA, ⁸³Department of Epidemiology and Population Health, Stanford University School of Medicine, Stanford, CA, USA, ⁸⁴Stanford Prevention Research Center, Department of Medicine, Stanford University, Stanford, CA, USA, ⁸⁵Meta-Research Innovation Center Berlin (METRIC-B), Berlin Institute of Health, Berlin, Germany

** Corresponding author:*

Lars G. Hemkens, MD, MPH

Department of Clinical Research, University Hospital Basel, Spitalstrasse 12, CH-4031 Basel, Switzerland

Phone: +41 61 265 3100, Mail: lars.hemkens@usb.ch

Word count: Main text: 2994. Abstract: 258.

Abstract

Substantial COVID-19 research investment has been allocated to randomized clinical trials (RCTs) on hydroxychloroquine/chloroquine, which currently face recruitment challenges or early discontinuation. We aimed to estimate the effects of hydroxychloroquine and chloroquine on survival in COVID-19 from all currently available RCT evidence, published and unpublished. We conducted a rapid meta-analysis of ongoing, completed, or discontinued RCTs on hydroxychloroquine or chloroquine treatment for any COVID-19 patients (protocol: <https://osf.io/QESV4/>). We systematically identified unpublished RCTs (ClinicalTrials.gov, WHO International Clinical Trials Registry Platform, Cochrane COVID-registry up to June 11, 2020), and published RCTs (PubMed, medRxiv and bioRxiv up to October 16, 2020). All-cause mortality was extracted (publications/preprints) or requested from investigators and combined in random-effects meta-analyses, calculating odds ratios (ORs) with 95% confidence intervals (CIs), separately for hydroxychloroquine and chloroquine. Prespecified subgroup analyses included patient setting, diagnostic confirmation, control type, and publication status. Sixty-three trials were potentially eligible. We included 14 unpublished trials (1308 patients) and 14 publications/preprints (9011 patients). Results for hydroxychloroquine are dominated by RECOVERY and WHO SOLIDARITY, two highly pragmatic trials, which employed relatively high doses and included 4716 and 1853 patients, respectively (67% of the total sample size). The combined OR on all-cause mortality for hydroxychloroquine was 1.11 (95% CI: 1.02, 1.20; $I^2=0\%$; 26 trials; 10,012 patients) and for chloroquine 1.77 (95% CI: 0.15, 21.13, $I^2=0\%$; 4 trials; 307 patients). We identified no subgroup effects. We found that treatment with hydroxychloroquine was associated with increased mortality in COVID-19 patients, and there was no benefit of chloroquine. Findings have unclear generalizability to outpatients, children, pregnant women, and people with comorbidities.

Keywords: Meta-analysis, SARS-CoV-2, COVID-19, Hydroxychloroquine, Chloroquine

Introduction

Coronavirus disease 2019 (COVID-19) caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) has the potential of progression into respiratory failure and death.¹ More than 900,000 persons with COVID-19 globally have died by September, 2020,² and treatment options are limited.³ The COVID-19 pandemic has caused a hitherto unprecedented search for possible therapies, with almost 700 clinical trials initiated in the first quarter of 2020 - and one in five of these trials target hydroxychloroquine (HCQ) or chloroquine (CQ).⁴ This remarkable attention was primarily due to *in vitro* data,⁵ immunomodulatory capacities,⁶ and the oral formulation and well-documented safety profiles.

In March 2020, the US Food and Drug Administration (FDA) issued an Emergency Use Authorization of HCQ⁷ and its prescription and usage outside clinical studies skyrocketed.⁸ In many countries, HCQ or CQ were listed in treatment guidelines for COVID-19 (including, e.g., China, Ireland, and the US).⁹ In a New York City cohort of 1376 COVID-19 inpatients during March-April 2020, 59% received HCQ.¹⁰ However, the FDA revoked the Emergency Use Authorization on June 15, 2020.¹¹ At that point, two large randomized clinical trials (RCTs), RECOVERY and the WHO Solidarity trial, had stopped enrollment to their HCQ treatment arms.^{12,13} An interim analysis of the RECOVERY trial showed no mortality benefit of HCQ.¹³ Established as treatments of malaria and rheumatic disorders, HCQ and CQ may carry potentially severe adverse effects, especially related to cardiac arrhythmia.⁶ Public uncertainty still remains, as illustrated by recent reports of planned use in pandemic epicenters in Central and South America.¹⁴

While many trials are ongoing, additional published evidence of potential benefits or harms may be several months away, if they even reach completion. Given the lack of favorable results in the large RECOVERY trial and the revoked Emergency Use Authorization, recruitment into HCQ and CQ trials has become increasingly difficult and many trials may run the risk of ending in futility. A rapid examination of data on all-cause mortality from as many trials as possible may offer the best evidence on potential survival benefits and to ensure that patients are not exposed to unnecessary risks if benefit is lacking. We used the infrastructure established with COVID-evidence,¹⁵ a comprehensive

database of COVID-19 trials funded by the Swiss National Science Foundation, to invite all investigators of HCQ or CQ trials to participate in an international collaborative meta-analysis. We aimed to identify and combine all RCTs investigating the effects of HCQ or CQ on all-cause mortality in patients with COVID-19 compared to any control arm similar to the experimental arm in all aspects except the administration of HCQ or CQ.

Methods

This collaborative meta-analysis focused solely on all-cause mortality in order to provide rapid evidence on the most critical clinical outcome. Investigators of ongoing, discontinued or completed trials were contacted via email to provide group-level (aggregated) mortality data per trial arm at any time point available. The protocol was published online before data collection.¹⁶

We considered all clinical trials that reported randomly allocating patients with confirmed or suspected SARS-CoV-2 infection to a treatment protocol containing HCQ or CQ (for any duration or dose) or the same treatment protocol not containing HCQ or CQ. In other words, the control group had to receive placebo or no treatment other than standard of care (we excluded comparisons of HCQ or CQ against an active treatment, e.g., HCQ versus azithromycin, since active controls were too heterogeneous to pool together and reveal the pure benefits and harms of HCQ or CQ). Eligible ongoing trials had to provide data on all-cause mortality and randomize the first patient before June 1, 2020 (time point selected arbitrarily as we did not expect trials launched later to recruit enough patients to provide relevant additional information). Trials published or posted as preprint were not restricted by date. Prevention trials were not included. We included trials regardless of whether mortality was a primary outcome or not and put no restrictions on trial status, language, geographical region, or healthcare setting.

We searched for eligible trials registered at ClinicalTrials.gov and the WHO International Clinical Trials Registry Platform [ICTRP] by June 11, 2020 (COVID-evidence database).¹⁷ We additionally searched PubMed and the Cochrane COVID-19 trial registry (covering preprints, trial registries and literature databases) by June 11, 2020, using terms related to HCQ and CQ combined with terms for COVID-19 and a standard RCT filter (Supplement 1).¹⁸ We updated the literature search on October 16, 2020. Two authors (CA and AMS) independently verified the eligibility criteria (Figure 1) and solved any discrepancies by discussion.

Principal investigators of 83 potentially eligible trials were asked to confirm the eligibility criteria, as well as: “For each of your study arms: (a) What intervention did this group receive? (b) How many patients were randomized to this group? (c) Of these patients, how many have died? (d) Of these patients, for how many it is unknown if they are dead or alive?” (Supplement 2, email template).

Investigators who were not responsive received two email reminders in English or Chinese, depending on trial origin.

The following information was extracted from all included RCTs by two reviewers (CA, AMS) and verified by the trial investigators: experimental and control arms, number of randomized participants, treatment schedule, patient setting, eligibility criteria, study location, blinding, target sample size, and trial status. We also classified trials as published in a peer-reviewed journal, posted on a preprint server, or unpublished (the latter category not including preprints). For reasons of feasibility within this rapid assessment, we generally did not request descriptive information beyond items included in trial registrations.

The main analysis evaluated separately the effect on all-cause mortality of HCQ versus control and CQ versus control. We report absolute numbers and proportions, as well as the treatment effect estimate as an odds ratio (OR; odds of death in the HCQ or CQ intervention group divided by the odds of death in the control group) with 95% confidence intervals (CIs). For multi-arm studies, we requested data for all arms and calculated treatment effect estimates for each eligible comparison. We combined mortality effects from all RCTs based on binary outcome data (2x2 contingency tables) in meta-analyses and describe the statistical heterogeneity using the I^2 -statistic.¹⁹ In our protocol, we prespecified a random-effects model of the Hartung-Knapp-Sidik-Jonkman (HKSJ) approach,²⁰ in order to provide more equality of weights between trials with moderate to large size (than for example the DerSimonian-Laird approach). We did not prespecify the between-study variance estimator, tau-squared, but chose the Paule and Mandel (PM) estimator based on provided guidance on choosing among 16 variants.²¹ Cases of zero events in one arm were corrected by adding the reciprocal of the size of the contrasting study arm.¹⁸ However, considering the range of sample sizes and numbers of zero events across trials, we assessed the effects of alternative approaches with sensitivity analyses, as

detailed below. To explore and illustrate evidence generation over time, we also performed a cumulative meta-analysis of all trials as well as stratified by dissemination status (publications/preprints vs unpublished), using the HKSJ approach with PM tau-squared. We used the date of email response or publication/posting of preprint. The meta-analyses were completed using R version 3.5.1 and the ‘meta’ package version 4.13-0.

In exploratory subgroup analyses, we stratified trials by patient setting (as proxy to COVID-19 severity: outpatients, inpatients but not intensive care unit (ICU), and ICU), diagnostic confirmation (confirmed SARS-CoV-2 versus suspected cases), control type (placebo control versus other) and publications/preprints versus unpublished trials. We did not stratify for missing data since the amount was extremely low. A post-hoc stratification by HCQ dose was added (trials with ≥ 1600 mg on day 1 and ≥ 800 mg from day 2 versus lower-dose trials) to isolate trials predicted to achieve blood levels of HCQ above the *in vitro* half maximal inhibitory concentration (IC50) value for SARS-CoV-2 (1.13 μM).²²

We added exploratory sensitivity analyses to assess robustness across meta-analytic approaches: DerSimonian-Laird and Sidik-Jonkman tau-squared estimators, Mantel-Haenszel random-effects method, and Peto method. DerSimonian-Laird is a standard random-effects meta-analysis approach, but may underestimate uncertainty. The Sidik-Jonkman tau-squared estimator, on the other hand, may yield inflated estimates if heterogeneity is low.²¹ The Mantel-Haenszel method performs reasonably well with small and zero event counts, much like Peto and arcsine transformation. The Peto method is suboptimal in the presence of substantial imbalances in the allocation ratio of patients randomized in the compared arms (e.g., RECOVERY trial). We also modeled variants to handling zero events (arcsine difference, and excluding trials with zero events) as well as excluding trials with <50 participants.

Results

Our search identified 146 randomized trials investigating HCQ or CQ as treatment for COVID-19, of which 83 were deemed potentially eligible after scrutinizing the randomized comparisons. The investigators of these 83 trials were contacted and 57% (47 of 83) responded (Figure 1). Of the responders, 19 trials were eligible and available (14 unpublished, three preprints and two publications); 21 trials were ineligible according to information provided; five responding investigator teams were not ready to share their results yet; and two declined participation. For the 36 trials without response, six were confirmed eligible and available (four publications and two preprints); two were confirmed ineligible; and for the remaining 28, results were not available, nor could they be confirmed eligible.

We included 28 trials (14 unpublished trials, eight publications, and six preprints; of these, one publication and two preprints were identified for the first time in our search update).^{13,23–35} Individual trial characteristics are presented in Table 1 (28 included trials) and Supplement Table S1 (35 potentially eligible but unavailable). Overall, trial characteristics were not different between included and unavailable trials (Table 2).

HCQ was evaluated in 26 trials (10,012 patients), and CQ was evaluated in four trials (307 patients). Two trials investigated both HCQ versus control and CQ versus control (63 patients). The median sample size was 95 (IQR 28 to 282) for HCQ trials and 42 (IQR 29 to 95) for CQ trials. The two largest trials (RECOVERY and WHO SOLIDARITY) included 47% and 19% of all patients in the HCQ trials, respectively. Most trials investigated HCQ or CQ in hospitalized patients (22 trials; 79%), and only five trials (18%) had an outpatient setting. The average mortality was 10.3% (standard deviation 13.5%) in inpatient trials and 0.08% (standard deviation 0.18%) in outpatient trials. The comparator was in eleven trials placebo (39%) and in 17 (61%) no other treatment than standard of care. In most trials, patients and clinicians were aware of the treatment (15 trials; 54%), while in one trial (4%) the patients were blinded and in 11 trials (39%) patients and clinicians were blinded (Table 2).

Regarding HCQ, in the 26 included trials, 606 of 4316 (14.0%) patients treated with HCQ died and 960 of 5696 patients (16.9%) in the control groups died (within 19 trials with a 1:1 randomization ratio, 7.7% of patients in the HCQ arm died [181 of 2346] and 7.1% of patients in the control arm died [168 of 2352]). In the meta-analysis, the combined OR was 1.11 (95% CI, 1.02 to 1.20, $p = 0.02$), with low heterogeneity ($I^2 = 0\%$) (Figure 2A). In 12 trials including a total of 1282 patients, there were zero deaths in both arms.

Regarding CQ, in the four included trials, 18 of 160 (11%) patients treated with CQ died and 12 of 147 patients (8%) in the control groups died. The combined OR was 1.77 (95% CI: 0.15 to 21.13, $p = 0.21$), with low heterogeneity ($I^2 = 0\%$) (Figure 2B). In two of four trials including a total of 217 patients, there were zero deaths in both arms.

The available evidence in this study is the result of publications, preprints or personal communication accrued from April 10, 2020 to October 15, 2020 as shown in the cumulative meta-analyses (Figure 3A-C).

For HCQ, none of the exploratory subgroup analyses showed an effect modification (Supplement Table S2A). When only including published information (publications and preprints, excluding unpublished trials), there was an increase in mortality among patients treated with HCQ (OR 1.12, 95% CI 1.08 to 1.16), while among the unpublished trials there was no such sign of increased mortality (OR 0.92, 95% CI 0.63 to 1.34, p for interaction = 0.23). We conducted no subgroup analyses for CQ, as there were only two trials with events. In the sensitivity analyses employing different meta-analytical approaches (Supplement Table S2B and Figures S1A-C), results were compatible with the main analysis.

Discussion

This collaborative meta-analysis of 28 published or unpublished RCTs, including 10,319 patients, shows that treatment with HCQ was associated with increased mortality in COVID-19 patients, and there was no benefit from treatment with CQ. No differences were seen across subgroup analyses on patient setting, diagnosis confirmation, control type, publication status or dose and the between-study heterogeneity was low. For CQ, the number of studies was too small to draw clear conclusions.

This meta-analysis offers useful insights for a challenging health situation. Hundreds of thousands of patients have received HCQ and CQ outside of clinical trials without evidence of their beneficial effects. Public interest is unprecedented, with weak early evidence supporting HCQ's merits being widely discussed in some media and social networks - despite the unfavorable results by a very large RCT. Numerous clinical studies have been investigating HCQ and CQ almost simultaneously.

Although several systematic reviews and meta-analyses are already available, they only consider the small handful of RCTs being already published (which were all included here).³⁶⁻⁴⁰ While data sharing has been rather limited to-date in biomedical research, such openness can be transformative in generating knowledge. This pandemic has brought together a collaboration of clinical trialists agreeing to share their data, which allows this study to not only summarize the existing evidence, but also illustrate the accumulation of evidence that would otherwise not be available.

For HCQ, evidence is dominated by the RECOVERY trial,¹³ which indicated no mortality benefit for treated COVID-19 patients, together with longer hospitalization and higher risk of progression to invasive mechanical ventilation and/or death. Similarly, the WHO SOLIDARITY trial indicated no mortality benefit.³⁵ The RECOVERY and WHO SOLIDARITY trials used HCQ in comparatively higher doses than all other trials except REMAP-CAP. There was no evidence for an effect modification by dose (p for interaction = 0.29), and the combined effect of all the trials with lower dose did not indicate a benefit of HCQ but tended to a null effect (OR 0.97; 0.73 to 1.30) with wide CIs, compatible with the main effect estimate.

Although the published trials resulted in a conclusive treatment estimate, the unpublished trials tend towards a null effect. The tendency of published trials to report larger effect sizes than unpublished trials is well-documented and constitute one of the reporting biases that are discernable only when a body of studies are considered together.⁴¹ Null results are less expected to be rapidly disseminated, especially if the trial is small. Of note, RECOVERY results showing dexamethasone benefits have been published more rapidly³ than the unfavorable HCQ results.¹³ This paper offers the most comprehensive summary on HCQ and mortality in COVID-19 to date.

This meta-analysis does not address prophylactic use nor other outcomes besides mortality. All but three trials excluded children and the majority excluded pregnant or breastfeeding women; generalizability remains unclear for those populations. Among the five studies on outpatients, there were three deaths, two occurring in the one trial of 491 relatively young patients with few comorbidities,²⁵ and one occurring in a small trial with 27 patients. For outpatients that are elderly or have comorbidities, evidence is sparse. Most of the 28 trials excluded persons with comorbid conditions carrying higher risk of adverse events from HCQ or CQ.²³⁻²⁵ No evidence is in the pipeline for these groups, which echoes clinical reasoning being reluctant to expose them to risk.

Twenty-three percent of the potentially eligible trials were listed as discontinued, mostly because of fewer patients than expected. Among 28 included RCTs, only two had reached their target sample size at the time of censoring for this meta-analysis. As previously discussed,⁴² most trials on HCQ and CQ in COVID-19 are small, reflecting both the strong motivation for individual efforts and underscoring the need for readily available research infrastructure to merge small-scale initiatives.⁴ Especially in the context of recruitment challenges, we encourage other researchers to form collaborations and combine trial results.⁴³

Our analysis has some limitations. First, although we adopted a comprehensive, systematic search strategy, our real-time initiative differs from traditional systematic reviews. We focused on collecting unpublished information, aiming to rapidly secure as much trial evidence as possible. We did not review individual trials, nor did we stratify results according to patient characteristics. Such analyses are planned in future publications using in-depth details disclosed in individual trial publications to

come.⁴⁴⁻⁴⁶ The exploratory subgroup analyses did not support the hypothesis that blinding/use of placebo is associated with the observed effect (the test for an interaction gives $p=0.15$ and the OR is 0.88 with wide CIs [0.55 to 1.41], compatible with the overall effect); moreover, attrition was negligible (median 0%, IQR 0% to 0%; range 0 to 19.5%). A meta-epidemiological study shows little evidence that mortality results would be affected by lack of blinding, or problems in randomization and allocation concealment, in contrast to less objective outcomes.⁴⁷ Second, a majority of the potentially eligible trials were not available. Despite going far beyond the standard review of published evidence, we expect additional results from future trials on CQ to narrow the uncertainty of the treatment effect and possibly reveal benefits or harms not discernible based on the current evidence. We plan to perform an update when substantial additional evidence becomes available. Finally, although sensitivity analyses addressing model specifications were compatible with the main analysis, one combination (HKSJ model with SJ tau-squared estimator) yielded substantially wider confidence intervals. This combination gave disproportionately low weight to RECOVERY (15%) and we consider the main model (HKSJ with PM tau-squared estimator) to be more valid in this situation.

Treatment with HCQ for COVID-19 was associated with increased mortality, and there was no benefit from CQ based on currently available randomized trial data. Medical professionals around the globe are encouraged to inform patients about this evidence.

Declarations

Availability of data and materials

All data generated or analysed during this study are included in this published article and its supplementary information files.

Competing interests

Benjamin Abella and Ravi Amaravadi are the primary investigators of the Prevention and Treatment of COVID19 with Hydroxychloroquine (PATCH) trial, funded by a philanthropic gift. Ravi Amaravadi reports being founder with equity of Pinpoint Therapeutics and Immunacell, and personal fees from Sprint Biosciences and Deciphera, outside the submitted work. Derek Angus reports personal fees from Ferring Pharmaceuticals, Inc., Bristol-Myers Squibb, and Bayer AG, other from Alung Technologies, Inc., outside the submitted work; in addition, Dr. Angus has pending patents for Selepressin - compounds, compositions and methods for treating sepsis to Ferring, B.V., and Proteomic biomarkers of sepsis in elderly patients pending to University of Pittsburgh. Yaseen Arabi reports that he is principal investigator on a clinical trial of lopinavir–ritonavir and interferon for Middle East respiratory syndrome (MERS) and that he was a non-paid consultant on therapeutics for MERS-coronavirus (CoV) for Gilead Sciences and SAB Biotherapeutics. He is a co-investigator on the Randomized, Embedded, Multi-factorial Adaptive Platform Trial for Community-Acquired Pneumonia (REMAP-CAP), a board member of the International Severe Acute Respiratory and Emerging Infection Consortium (ISARIC), and the Lead-Co Chair of the Think20 (T20) Taskforce for COVID-19. Brigham and Women’s Hospital, PRA Health Science, and Cliniques universitaires Saint-Luc received funds from Sanofi. Thomas Benfield reports grants from Pfizer, Novo Nordisk Foundation, Simonsen Foundation, Lundbeck Foundation, and Kai Hansen Foundation; grants and personal fees from GSK, Pfizer, Boehringer Ingelheim, and Gilead; and personal fees from MSD, all outside the submitted work. Yehuda Cohen, Lisa Cowan, Bruno Igau, and Laurent Perrin are employees of Sanofi. The COV-HCQ and COMIHY trials were supported by the German Federal Ministry of Education and Research (EudraCT number 2020-001224-33) and the German Federal

Ministry of Health (EudraCT number 2020-001512-26). Lennie Derde reports grants from EU FP7-HEALTH-2013-INNOVATION-1, grant number 602525, grants from H2020 RECOVER grant agreement No 101003589, during the conduct of the study; and is a member of the COVID-19 guideline committee SCCM/ESICM/SSC, member of the ESICM COVID-19 taskforce, and chair of the Dutch intensivists (NVIC) taskforce infectious threats. Vincent Dubee reports non-financial support from MSD France and from Sanofi Aventis France, outside the submitted work. Anissa Elfakir is an employee of Ividata Life Sciences and works as an external contractor for Sanofi. Anthony Gordon received grant funding from a NIHR Research Professorship (RP-2015-06-18), support from the NIHR Imperial Biomedical Research Centre, and consulting fees paid to his institution from GlaxoSmithKline and Bristol Myers Squibb. Thomas Hills reports grants from the Health Research Council of New Zealand, during the conduct of the study. Andy Hoepelman reports grants from ZonMw, Netherlands organisation for Health Research and Development, during the conduct of the study. HYDRA trial was an investigator-initiated study supported by Sanofi, CONACYT (National Council of Science and Technology of Mexico) and by the participating centers. Thuy Le reports grants from Gilead Sciences, outside the submitted work. Bryan McVerry reports grants from NIH/NHLBI, and from Bayer Pharmaceuticals, Inc., outside the submitted work. Srinivas Murthy receives funding as the Innovative Medicines Canada Chair in Pandemic Preparedness. Colin McArthur reports grants from the Health Research Council of New Zealand. Mark Mulligan reports having received the HCQ drug from the New York State government, during the conduct of the study; grants from Lilly, Pfizer, Sanofi, and personal fees from Meissa, outside the submitted work; in addition, Dr. Mulligan has a patent anti-Zika monoclonal ab/ Emory Univ pending. Alistair Nichol is supported by a Health Research Board of Ireland Clinical Trial Network Award (HRB-CTH-2014-012). Lena Perez is an employee of Excelya and works as an external contractor for Sanofi. Frank Rockhold reports personal fees from Merck Research Labs, Novartis, Lilly, Sanofi, NovoNordisk, KLSMC, Tolerion, Rhythm, UCB, AstraZeneca, Janssen, Merck KGaA, Sarepta, Eidos, Amgen, Phathom, outside the submitted work; and having equity interest in GlaxoSmithkline, Athira Pharma, DataVant, Spencer Healthcare. Stephen Walsh reports receiving a grant from Sanofi during the conduct of the study and grants from NIH-NIAID outside the submitted

work, and having conducted vaccine (HIV, Zika) clinical trials funded by Janssen. Steve Webb reports grants from National Health and Medical Research Council (Australia), grants from Minderoo Foundation, from Health Research Council (New Zealand), and from Medical Research Future Fund (Australia), during the conduct of the study. Jesper Weehuizen reports grants from ZonMw, Netherlands organisation for Health Research and Development, during the conduct of the study. Fernando Zampieri was part of the Coalition 1 trial partially supported by EMS Pharmaceuticals, has received previous grants from Bactiguard, Sweden, outside the submitted work and support from Baxter LA for another clinical trial in critically ill patients. None of the other authors have any competing interests to declare.

Funding

This collaborative meta-analysis was supported by the Swiss National Science Foundation and Laura and John Arnold Foundation (grant supporting the post-doctoral fellowship at the Meta-Research Innovation Center at Stanford (METRICS), Stanford University). Funding also includes postdoctoral grants from Uppsala University, the Swedish Society of Medicine, the Blanceflor Foundation and the Sweden-America Foundation (C. Axfors). The funders had no role in the design of this collaborative meta-analysis; in the collection, analysis, and interpretation of data; or in the report writing. The corresponding author had full access to all study data and final responsibility for the decision to submit for publication.

Authors' contributions

Lars G. Hemkens, Cathrine Axfors and Andreas M. Schmitt had full access to all data in this study and take responsibility for the integrity of the data and the accuracy of the data analysis.

Concept and design: Lars G. Hemkens, John P. A. Ioannidis, Cathrine Axfors, Andreas M. Schmitt, Steven N. Goodman, David Moher

Acquisition, analysis, or interpretation of data: All authors

Drafting of the manuscript: Cathrine Axfors, Andreas M. Schmitt, Lars G. Hemkens

Critical revision of the manuscript for important intellectual content: All authors

Statistical analysis: Andreas M. Schmitt, Lars G. Hemkens, John P. A. Ioannidis, Steven N.

Goodman, Perrine Janiaud

Approval of the final manuscript: All authors

Obtained funding: Lars G. Hemkens, Cathrine Axfors, John P. A. Ioannidis

Administrative, technical, or material support: Cathrine Axfors, Andreas M. Schmitt, Lars G.

Hemkens, Perrine Janiaud, Janneke van 't Hooft

Supervision: Lars G. Hemkens

Acknowledgements

We wish to express our heartfelt gratitude to all patients volunteering for the trials involved. We furthermore thank Wenyan Ma and Benjamin Kasenda (University Hospital Basel, University of Basel) for kindly translating the emails to Chinese investigators. For valuable contributions to individual trials in this collaborative group, we sincerely thank: Hannah Jin, Monica Feeley, Bruce Bausk, Jessica Cauley, Jane Kleinjan, Jon Gothing, Naeema Bangash, Heather Wroe, Claire Bigogne, Christelle Castell, Annelies Mottart, Lisette Cortenraad, Judith Medema-Muller, Katia Handelberg, Khalid Benhammou, Shaheen Kumar, Sophie Gribomont, Kim Kuehne, Cathia Markina, Julien Labeirie, Julie Pencole, Eva Crispyn, Cecile Le Breton, Kelli Horn, Tina Patel, Benjamin Harrois, Isabelle Collin, Vetheeswar Manivannan, Irma Slomp, Frederick Becue, Isabelle Godefroy, Lynne Guo, Lene Kollmorgen, Toluwalope Cole, Catherine Chene, Praveena Deenumsetti, Anne Doisy, Ariane Vialfont, Melissa Charbit, Christine Shu, Stephane Kirkesseli, Howard Surks, Magalie De Meyer, Edel Hendrickx, and Paul Deutsch (Sanofi trial); Ellie Carmody, Märtin Backer, Jaishvi Eapen, Jack A. DeHovitz, Prithiv J. Prasad, Yi Li, Camila Delgado, Morris Jrada, Gabriel A. Robbins, Brooklyn Henderson, Alexander Hrycko, Dinuli Delpachitra, Vanessa Raabe, Jonathan S. Austrian, and Yanina Dubrovskaya (TEACH trial); Farah Al-Beidh, Djillali Annane, Kenneth Baillie, Abigail Beane, Richard Beasley, Zahra Bhimani, Marc Bonten, Charlotte Bradbury, Frank Brunkhorst, Meredith Buxton, Allen Cheng, Menno de Jong, Eamon Duffy, Lise Estcourt, Rob Fowler, Timothy Girard, Herman Goossens, Cameron Green, Rashan Haniffa, Christopher Horvat, David Huang, Francois Lamontagne, Patrick Lawler, Kelsey Linstrum, Edward Litton, John Marshall, Daniel

McAuley, Shay McGuinness, Stephanie Montgomery, Paul Mouncey, Katrina Orr, Rachael Parke, Jane Parker, Asad Patanwala, Kathryn Rowan, Marlene Santos, Christopher Seymour, Steven Tong, Anne Turner, Timothy Uyeki, Wilma van Bentum-Puijk, Frank van de Veerdonk, and Ryan Zarychanski (REMAP-CAP trial); Jan-Erik Berdal, Arne Eskesen, Dag Kvale, Inge Christoffer Olsen, Corina Silvia Rueegg, Anbjørg Rangberg, Christine Monceyron Jonassen, and Torbjørn Omland (NO COVID-19 Trial). Support for title page creation and format was provided by AuthorArranger, a tool developed at the National Cancer Institute.

References

1. Berlin, D. A., Gulick, R. M. & Martinez, F. J. Severe Covid-19. *N. Engl. J. Med.* (2020)
doi:10.1056/NEJMcp2009575.
2. COVID-19 Map. *Johns Hopkins Coronavirus Resource Center*
<https://coronavirus.jhu.edu/map.html>.
3. RECOVERY Collaborative Group *et al.* Dexamethasone in Hospitalized Patients with Covid-19 - Preliminary Report. *N. Engl. J. Med.* (2020) doi:10.1056/NEJMoa2021436.
4. Janiaud, P. *et al.* The worldwide clinical trial research response to the COVID-19 pandemic - the first 100 days. *Fl1000Res.* **9**, 1193 (2020).
5. Fantini, J., Di Scala, C., Chahinian, H. & Yahi, N. Structural and molecular modelling studies reveal a new mechanism of action of chloroquine and hydroxychloroquine against SARS-CoV-2 infection. *Int. J. Antimicrob. Agents* **55**, 105960 (2020).
6. dos Reis Neto, E. T. *et al.* Revisiting hydroxychloroquine and chloroquine for patients with chronic immunity-mediated inflammatory rheumatic diseases. *Advances in Rheumatology* **60**, 32 (2020).
7. Office of the Commissioner. Emergency Use Authorization. *U.S. Food and Drug Administration*
<https://www.fda.gov/emergency-preparedness-and-response/mcm-legal-regulatory-and-policy-framework/emergency-use-authorization> (2020).
8. Vaduganathan, M. *et al.* Prescription Fill Patterns for Commonly Used Drugs During the COVID-19 Pandemic in the United States. *JAMA* (2020) doi:10.1001/jama.2020.9184.
9. Dagens, A. *et al.* Scope, quality, and inclusivity of clinical guidelines produced early in the covid-19 pandemic: rapid review. *BMJ* **369**, m1936 (2020).
10. Geleris, J. *et al.* Observational Study of Hydroxychloroquine in Hospitalized Patients with Covid-19. *N. Engl. J. Med.* (2020) doi:10.1056/NEJMoa2012410.
11. Office of the Commissioner. Coronavirus (COVID-19) Update: FDA Revokes Emergency Use Authorization for Chloroquine and Hydroxychloroquine. *U.S. Food and Drug Administration*
<https://www.fda.gov/news-events/press-announcements/coronavirus-covid-19-update-fda>

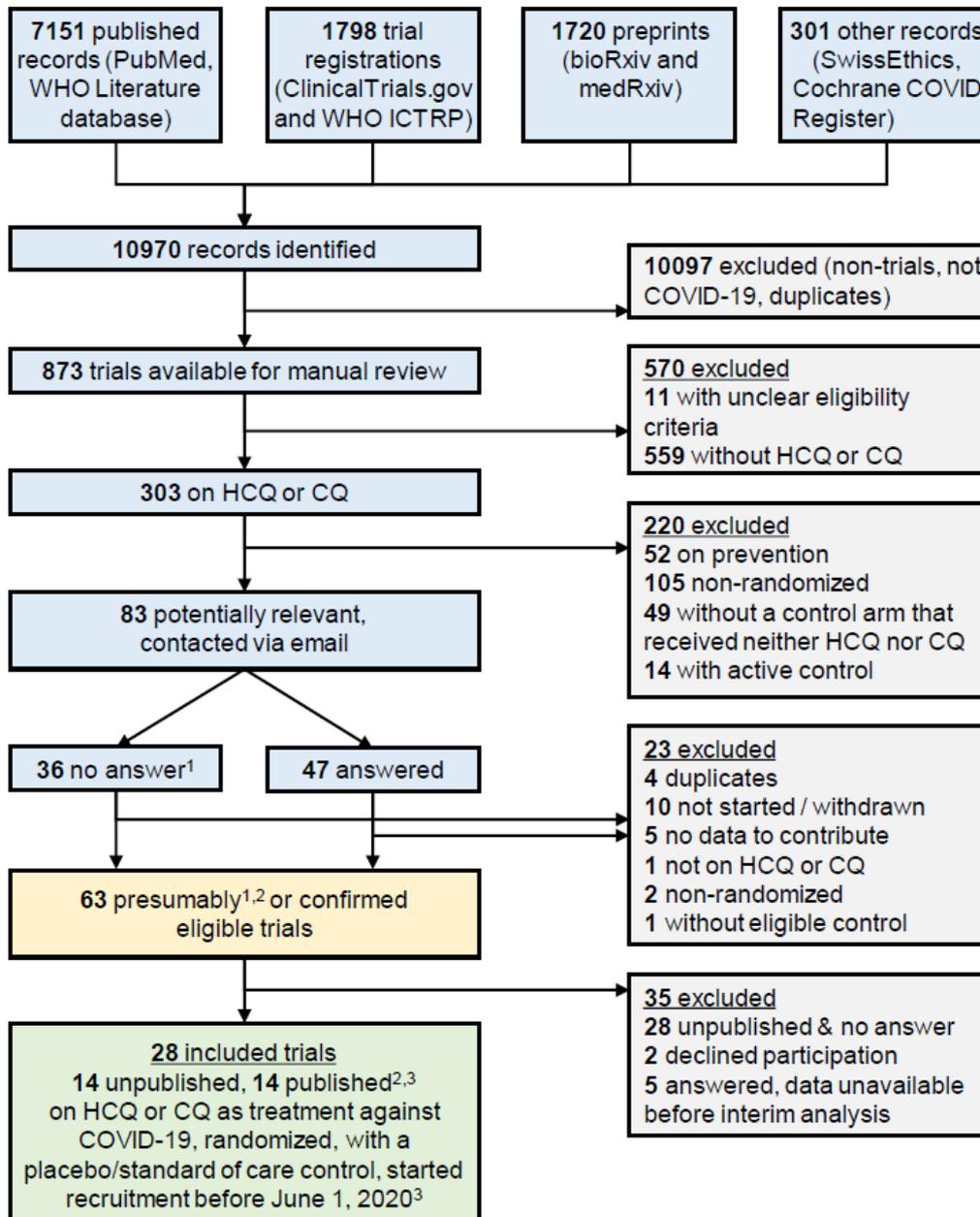
- revokes-emergency-use-authorization-chloroquine-and (2020).
12. WHO discontinues hydroxychloroquine and lopinavir/ritonavir treatment arms for COVID-19. <https://www.who.int/news-room/detail/04-07-2020-who-discontinues-hydroxychloroquine-and-lopinavir-ritonavir-treatment-arms-for-covid-19>.
 13. RECOVERY Collaborative Group *et al.* Effect of Hydroxychloroquine in Hospitalized Patients with Covid-19. *N. Engl. J. Med.* (2020) doi:10.1056/NEJMoa2022926.
 14. Trigo, M. S., Kurmanaev, A. & Cabrera, J. M. L. With Officials' Backing, Dubious Virus Remedies Surge in Latin America. *The New York Times* (2020).
 15. Janiaud, P., Axfors, C., Sacchilotto, R. & Hemkens, L. COVID-evidence: a living database of trials on interventions for COVID-19. (2020) doi:10.17605/OSF.IO/GEHFX.
 16. Axfors, C. *et al.* Hydroxychloroquine and chloroquine for survival in COVID-19: an international collaborative meta-analysis of randomized trials. (2020) doi:10.17605/OSF.IO/QESV4.
 17. COVID-evidence. <https://covid-evidence.org/>.
 18. Higgins, J. P. T. *et al.* *Cochrane Handbook for Systematic Reviews of Interventions*. (John Wiley & Sons, 2019).
 19. Higgins, J. P. T., Thompson, S. G., Deeks, J. J. & Altman, D. G. Measuring inconsistency in meta-analyses. *BMJ* **327**, 557–560 (2003).
 20. Int'Hout, J., Ioannidis, J. P. A. & Borm, G. F. The Hartung-Knapp-Sidik-Jonkman method for random effects meta-analysis is straightforward and considerably outperforms the standard DerSimonian-Laird method. *BMC Med. Res. Methodol.* **14**, 25 (2014).
 21. Veroniki, A. A. *et al.* Methods to estimate the between-study variance and its uncertainty in meta-analysis. *Res Synth Methods* **7**, 55–79 (2016).
 22. White, N. J. *et al.* COVID-19 prevention and treatment: a critical analysis of chloroquine and hydroxychloroquine clinical pharmacology. (2020).
 23. Chen, L. *et al.* Efficacy and safety of chloroquine or hydroxychloroquine in moderate type of COVID-19: a prospective open-label randomized controlled study. *Infectious Diseases (except HIV/AIDS)* (2020) doi:10.1101/2020.06.19.20136093.

24. Tang, W. *et al.* Hydroxychloroquine in patients with mainly mild to moderate coronavirus disease 2019: open label, randomised controlled trial. *BMJ* **369**, m1849 (2020).
25. Skipper, C. P. *et al.* Hydroxychloroquine in Nonhospitalized Adults With Early COVID-19: A Randomized Trial. *Ann. Intern. Med.* (2020) doi:10.7326/M20-4207.
26. Jun, C. *et al.* A pilot study of hydroxychloroquine in treatment of patients with moderate COVID-19. *J. Zhejiang Univ.* **49**, 215–219 (2020).
27. Cavalcanti, A. B. *et al.* Hydroxychloroquine with or without Azithromycin in Mild-to-Moderate Covid-19. *N. Engl. J. Med.* (2020) doi:10.1056/NEJMoa2019014.
28. Mitjà, O. *et al.* Hydroxychloroquine for Early Treatment of Adults with Mild Covid-19: A Randomized-Controlled Trial. *Clin. Infect. Dis.* (2020) doi:10.1093/cid/ciaa1009.
29. Chen, Z. *et al.* Efficacy of hydroxychloroquine in patients with COVID-19: results of a randomized clinical trial. *medRxiv* 2020.03.22.20040758 (2020).
30. Lyngbakken, M. N. *et al.* A pragmatic randomized controlled trial reports the efficacy of hydroxychloroquine on coronavirus disease 2019 viral kinetics. *In Review* (2020) doi:10.21203/rs.3.rs-44055/v1.
31. Chen, C.-P. *et al.* A Multicenter, randomized, open-label, controlled trial to evaluate the efficacy and tolerability of hydroxychloroquine and a retrospective study in adult patients with mild to moderate Coronavirus disease 2019 (COVID-19). *medRxiv* 2020.07.08.20148841 (2020).
32. Kamran, S. M. *et al.* Clearing the fog: Is HCQ effective in reducing COVID-19 progression: A randomized controlled trial. *medRxiv* 2020.07.30.20165365 (2020).
33. Ulrich, R. J. *et al.* Treating Covid-19 With Hydroxychloroquine (TEACH): A Multicenter, Double-Blind, Randomized Controlled Trial in Hospitalized Patients. *Open Forum Infect Dis* doi:10.1093/ofid/ofaa446.
34. Abd-Elsalam, S. *et al.* Hydroxychloroquine in the Treatment of COVID-19: A Multicenter Randomized Controlled Study. *Am. J. Trop. Med. Hyg.* (2020) doi:10.4269/ajtmh.20-0873.
35. WHO Solidarity Trial Consortium *et al.* Repurposed antiviral drugs for COVID-19; interim WHO SOLIDARITY trial results. *medRxiv* 2020.10.15.20209817 (2020).
36. Siemieniuk, R. A. *et al.* Drug treatments for covid-19: living systematic review and network

- meta-analysis. *BMJ* **370**, m2980 (2020).
37. Thoguluva Chandrasekar, V. *et al.* Systematic review and meta-analysis of effectiveness of treatment options against SARS-CoV-2 infection. *J. Med. Virol.* (2020) doi:10.1002/jmv.26302.
 38. Liu, W. *et al.* Efficacy and safety of antiviral treatment for COVID-19 from evidence in studies of SARS-CoV-2 and other acute viral infections: a systematic review and meta-analysis. *CMAJ* **192**, E734–E744 (2020).
 39. Hernandez, A. V., Roman, Y. M., Pasupuleti, V., Barboza, J. J. & Michael White, C. Hydroxychloroquine or Chloroquine for Treatment or Prophylaxis of COVID-19: A Living Systematic Review. *Ann. Intern. Med.*
 40. Sarma, P. *et al.* Virological and clinical cure in COVID-19 patients treated with hydroxychloroquine: A systematic review and meta-analysis. *J. Med. Virol.* **92**, 776–785 (2020).
 41. Dechartres, A., Atal, I., Riveros, C., Meerpohl, J. & Ravaud, P. Association Between Publication Characteristics and Treatment Effect Estimates: A Meta-epidemiologic Study. *Ann. Intern. Med.* **169**, 385–393 (2018).
 42. Janiaud, P. *et al.* 100 days later - the worldwide clinical trial research agenda of the COVID-19 pandemic.
 43. Petkova, E., Antman, E. M. & Troxel, A. B. Pooling Data From Individual Clinical Trials in the COVID-19 Era. *JAMA* (2020) doi:10.1001/jama.2020.13042.
 44. Bravo-Jeria, R. *et al.* Chloroquine and hydroxychloroquine for the treatment of COVID-19: a living systematic review. *PROSPERO 2020 CRD42020178195* (2020).
 45. Scott, D. A. *et al.* A systematic review and meta-analysis of the efficacy and safety of chloroquine for the treatment of patients with COVID-19. *PROSPERO 2020 CRD42020178266* (2020).
 46. Fontes, L. E. *et al.* Chloroquine/hydroxychloroquine for coronavirus disease 2019 (COVID-19) – a systematic review of individual participant data. *PROSPERO 2020 CRD42020178667*.
 47. Savović, J. *et al.* Influence of reported study design characteristics on intervention effect estimates from randomized, controlled trials. *Ann. Intern. Med.* **157**, 429–438 (2012).

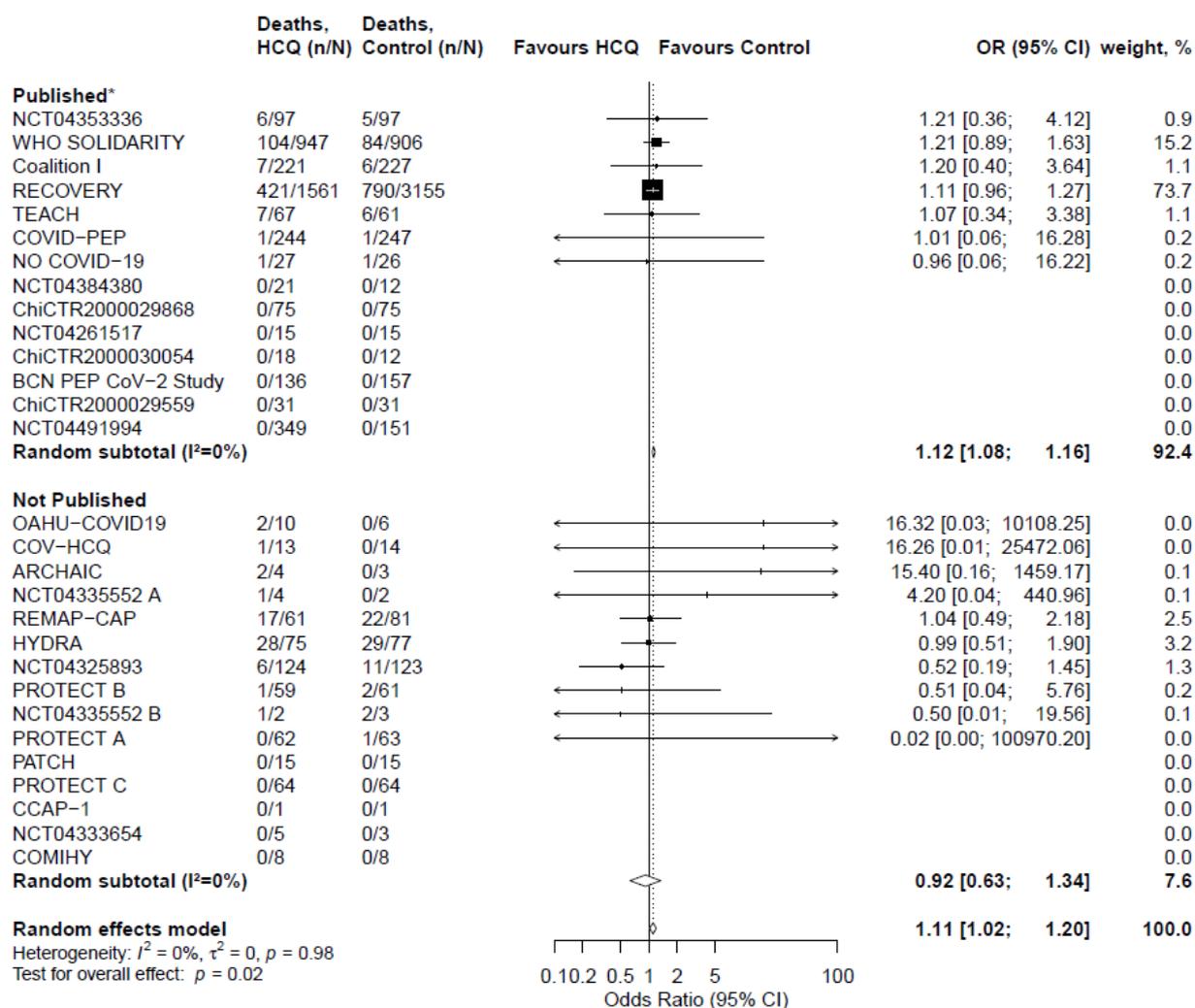
Figure captions

Figure 1. Flowchart of included randomized clinical trials.



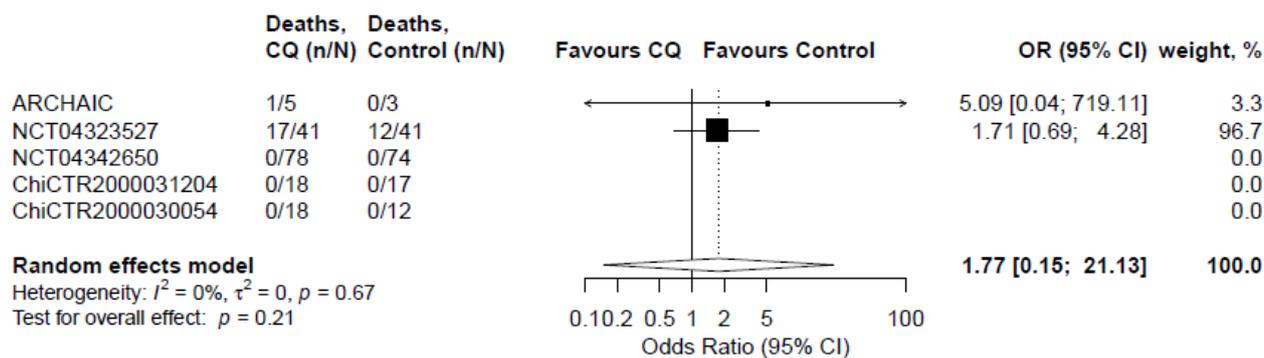
Sources searched up to June 11, 2020 (PubMed, ClinicalTrials.gov, WHO ICTRP, Cochrane COVID Register) or April 9 (WHO Literature database, bioRxiv, medRxiv, SwissEthics). ¹ Trials for which we received no answer were presumed to be eligible unless withdrawn. ² One publication and two preprints were identified in a later search update. ³ Published peer-reviewed articles or posted preprints. Abbreviations: chloroquine (CQ), hydroxychloroquine (HCQ), International Clinical Trials Registry Platform (ICTRP), World Health Organization (WHO).

Figure 2A. Random effects meta-analysis for mortality for treatment of COVID-19 with Hydroxychloroquine, trials are stratified by publication status.



* Published as peer-reviewed articles or posted preprints.

Figure 2B. Random effects meta-analysis for mortality for treatment of COVID-19 with Chloroquine.



The x-axis scales differ for reasons of readability.

Figure 3A. Cumulative meta-analysis for mortality for treatment of COVID-19 with Hydroxychloroquine.

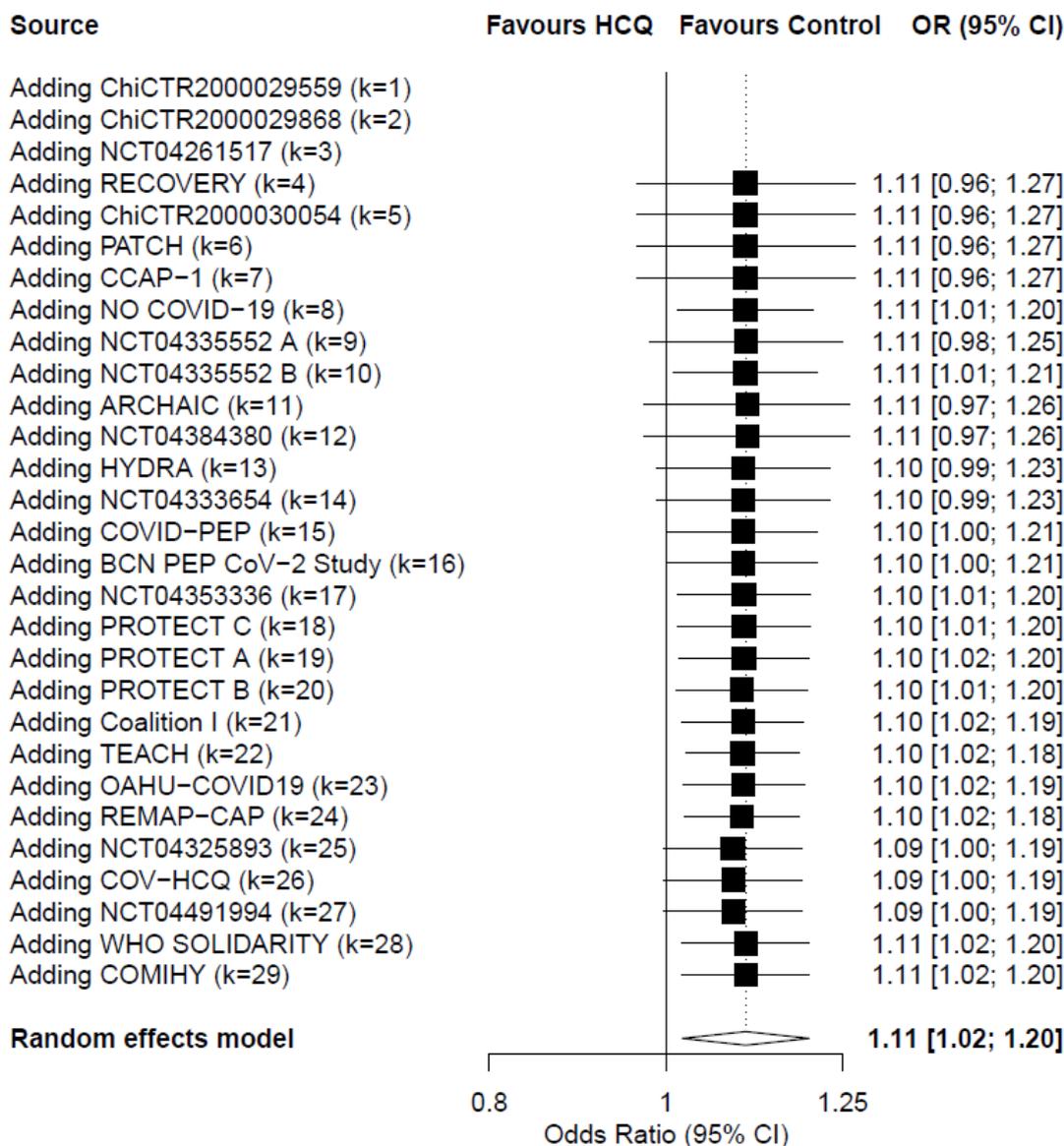


Figure 3B. Cumulative meta-analysis for mortality for treatment of COVID-19 with Hydroxychloroquine (publications and preprints only)

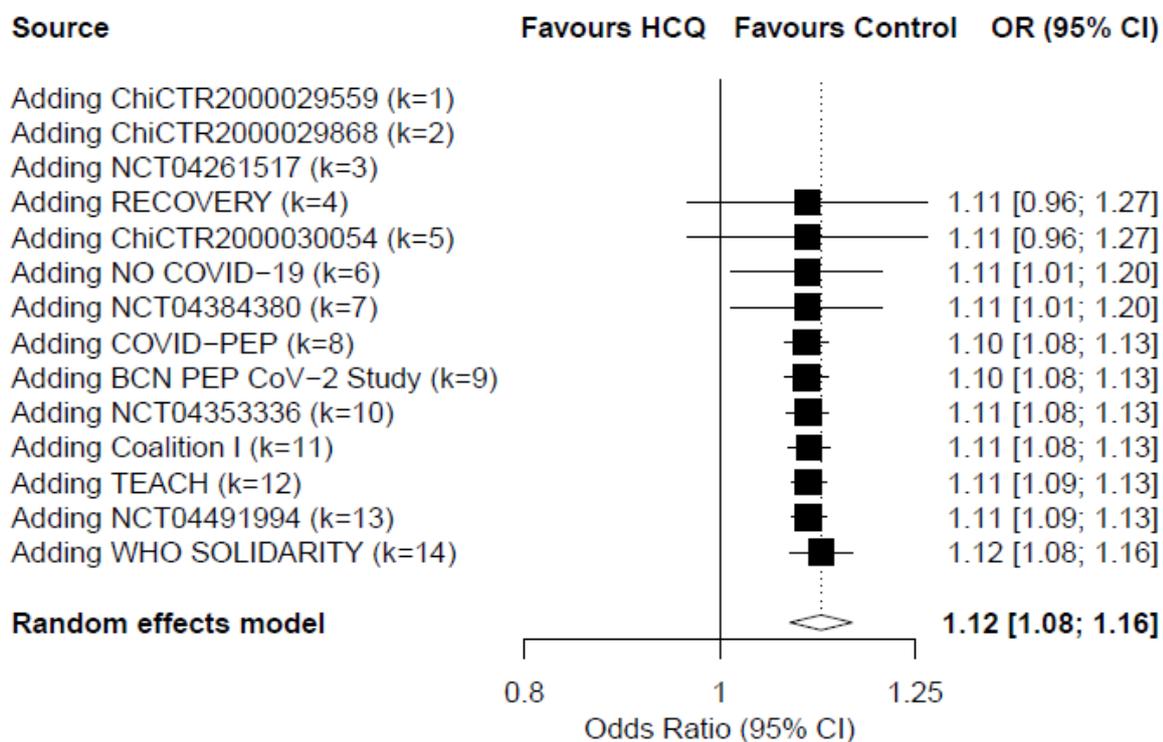
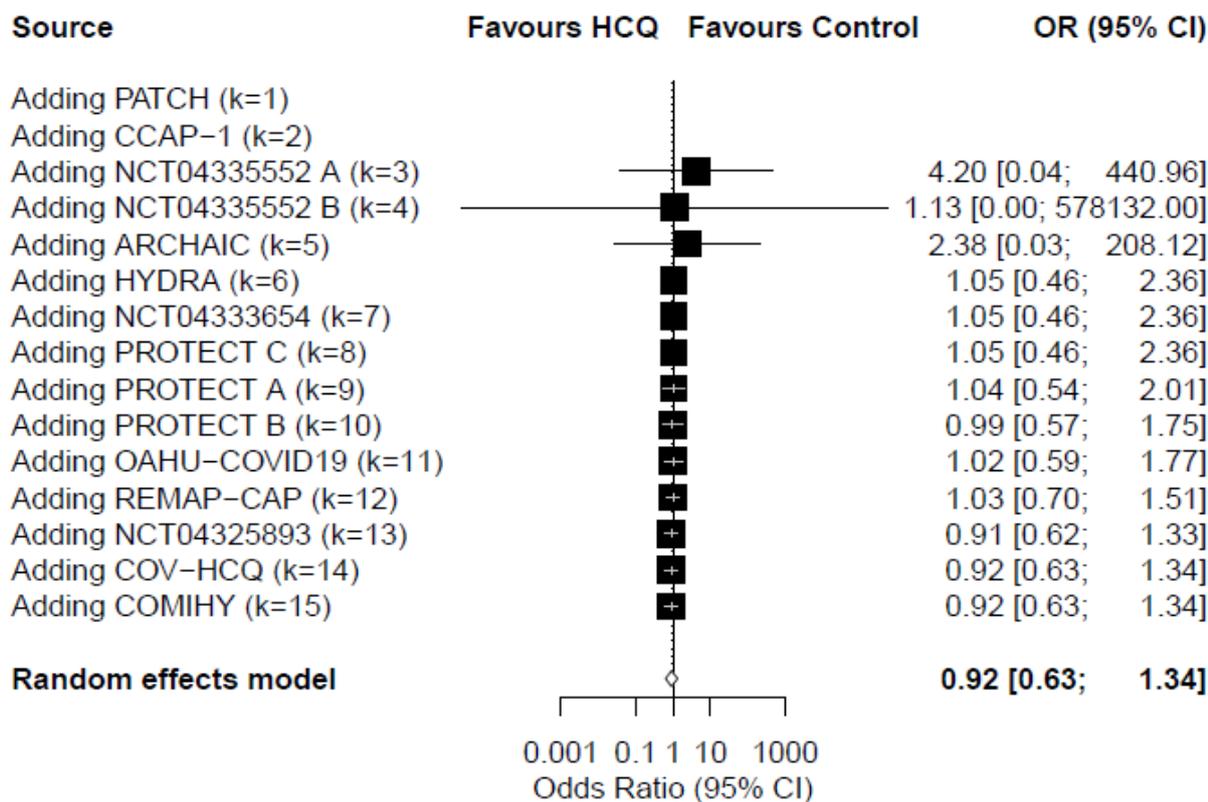


Figure 3C. Cumulative meta-analysis for mortality for treatment of COVID-19 with Hydroxychloroquine
(unpublished data only)



The x-axis scales differ for reasons of readability.

Table 1. Group-level characteristics of randomized clinical trials evaluating hydroxychloroquine or chloroquine as treatment for COVID-19.

Acronym	Register ID	Treatment comparison		Treatment schedule	Setting	Age	Mortality (%)	Location	Blinding	Targeted sample size	Status
		Experimental arm (n)	Control arm(n)								
<i>Published****</i>											
-	ChiCTR2000029559	HCQ (31)	No Treatment (31)	200 mg twice a day for 5 days	Inpatient	≥18 years	0	China	Participant, Caregiver	300	Completed
-	ChiCTR2000029868	HCQ (75)	No Treatment (75)	1200 mg/day for 3 days, then 800 mg/day for 11-18 days	Inpatient	≥18 years	0	China	None	360	Completed
-	NCT04261517	HCQ (15)	No Treatment (15)	400 mg/day for 5 days	Inpatient	≥18 years	0	China	None	30	Completed
RECOVERY	NCT04381936	HCQ (1561)	No Treatment (3155)	800 mg at zero hours, then 800 mg after 6 hours, then 800 mg/day for up to 9 days	Inpatient	≥18 years	25.57	United Kingdom	None	12000*	Completed **
-	ChiCTR2000030054	HCQ (18)	No Treatment (12)	400 mg/day for 10 days	Inpatient	18 to 75 years	0	China	None	100	Completed
		CQ (18)	No Treatment (12)	1000 mg/day for 1 day, then 500 mg/days for 9 days			0				
NO COVID-19	NCT04316377	HCQ (27)	No Treatment (26)	800 mg/day for 7 days	Inpatient	≥18 years	3.77	Norway	None	202	Halted
-	NCT04384380	HCQ (21)	No Treatment (12)	800 mg/day for 1 day, then 400 mg/day for 6 days	Inpatient	20 to 79 years	0	Taiwan	None	45	Recruiting
COVID-PEP	NCT04308668	HCQ (244)	Placebo (247)	800 mg at zero hours, then 600 mg after 6-8 hours, then 600 mg daily for 4 days	Outpatient	≥18 years	0.41	International ***	Participant, Caregiver	3000	Completed
BCN PEP CoV-2	NCT04304053	HCQ (136)	No Treatment (157)	800mg on day 1, and 400mg/day on days 2-7	Outpatient	≥18 years	0	Spain	None	2300	Completed
-	NCT04353336	HCQ (97)	No Treatment (97)	800 mg/day on day 1, then 400 mg/day for 14 days	Inpatient	All	5.67	Egypt	None	40	Recruiting
Coalition I	NCT04322123	HCQ (221)	No Treatment (227)	800 mg/day for 7 days	Inpatient	≥18 years	2.90	Brazil	None	630*	Completed
TEACH	NCT04369742	HCQ (67)	Placebo (61)	800 mg/day on day 1, then 400 mg/day for 4 days	Inpatient	All	10.16	United States	Participant, Caregiver	626	Discont.

	NCT04491994	HCQ (349)	No Treatment (151)	400 mg twice a day on day1, then 200 mg twice a day for 4 days	Inpatient	All	0	Pakistan	None	540	Completed
WHO SOLIDARITY	2020-001366-11	HCQ (947)	No Treatment (906)	Day 1, 2000 mg (Hour 0, 800 mg; Hour 6, 800 mg; Hour 12, 400 mg), thereafter 800 mg per day for 10 days	Inpatient	≥18 years	10.15%	International	None	10000*	Discont.
Unpublished											
PATCH	NCT04329923	HCQ (15)	Placebo (15)	800 mg/day for up to 14 days	Inpatient	≥40 years	0	United States	Participant, Caregiver	400*	Recruiting
CCAP-1	NCT04345289	HCQ (1)	Placebo (1)	600 mg/day for 7 days	Inpatient	≥18 years	0	Denmark	Participant, Caregiver	1500*	Discont.
-	NCT04335552	HCQ (4) HCQ + Azithromycin (2)	No Treatment (2) Azithromycin (3)	800 mg/day for 1 day, then 600 mg/day for 4 days	Inpatient	≥12 years	16.67 60	United States	None	500	Discont.
ARCHAIC	NL8490	CQ (5) HCQ (4)	No Treatment (3) No Treatment (3)	600 mg at zero hours, then 300 mg after 12 hours, then 600 mg/day for 4 days 800 mg/day for 1 day, then 400 mg/day for 4 days	Inpatient	≥18 years	12.50 28.57	Netherlands	None	950	Discont.
CloroCOVID19II	NCT04342650	CQ (78)	Placebo (74)	900 mg/day for 1 day, then 450 mg/day for 4 days	Outpatient	≥18 years	0	Brazil	Participant, Caregiver	210	Completed
	NCT04323527	CQ (41)	Placebo (41)	900 mg/day for 1 day, then 450 mg/day for 4 days	Inpatient	≥18 years	35.37	Brazil	Participant, Caregiver	278	Completed
HYDRA	NCT04315896	HCQ (75)	Placebo (77)	400 mg/day for 10 days	Inpatient	18 to 80 years	37.50	Mexico	Participant, Caregiver	500	Recruiting
-	ChiCTR2000031204	CQ (18)	Placebo (17)	1000 mg on day 1, then 500 mg/day on days 2-3, then 250 mg/day until ≤14 days of total treatment	Inpatient	18 to 70 years	0	China	Participant	300	Recruiting
-	NCT04333654	HCQ (5)	Placebo (3)	800 mg at zero hours, then 400 mg 6-8 hours later, then 600 mg/day for 9 days	Outpatient	18 to 80 years	0	International ***	Participant, Caregiver	210	Discontin
PROTECT	NCT04338698	HCQ + Azithromycin + Oseltamivir (64) HCQ + Oseltamivir (62)	Azithromycin + Oseltamivir (64) Oseltamivir (63)	600 mg/day for 5 days	Inpatient	≥18 years	0 0.80	Pakistan	Investigator	500	Recruiting

		HCQ + Azithromycin (59)	Azithromycin (61)				2.50				
OAHU-COVID19	NCT04345692	HCQ (10)	No Treatment (6)	800 mg/day on day 1, then 400 mg/day for 4 days	Inpatient	18 to 95 years	12.50	United States	None	350	Recruiting
REMAP-CAP	NCT02735707	HCQ (61)	No Treatment (81)	800 mg at zero and six hours, then 800 mg/day for up to 6 days	ICU	≥18 years	27.46	International ***	None	No fixed target sample	Completed **
-	NCT04325893	HCQ (124)	Placebo (123)	800 mg on day 1, then 400 mg/day for 8 days	Inpatient	≥18 years	6.88	France	Participant, Caregiver	1300	Recruiting
COV-HCQ*****	NCT04342221	HCQ (13)	Placebo (14)	800mg/day on day 1 and 600 mg/day for days 2-7	Inpatient	≥18 years	3.70	Germany	Participant, Caregiver	220	Recruiting
COMIHY	NCT04340544	HCQ (8)	Placebo (8)	600mg/day for 7 days	Outpatient	≥18 years	0	Germany	Participant, Caregiver	2700	Recruiting

Abbreviations: chloroquine (CQ), hydroxychloroquine (HCQ). * Trial includes more treatment arms than reported here; target sample size refers to all arms. **

Other arms of the trial are still ongoing. *** Including centers in multiple countries. **** Including peer-reviewed journal publications and posted preprints.

***** The allocation of participants to the experimental arm (n = 13) and control arm (n = 14) versus experimental arm (n = 14) and control arm (n = 13) was not yet unblinded. The allocation for the meta-analysis was done randomly. A sensitivity analysis with the reverse assignment gave the same result.

Table 2. Group-level characteristics of included and unavailable trials.

	All trials n = 63	Included trials n = 28	Potentially eligible, unavailable trials* n = 35
Drug, n (%)			
HCQ	48 (76)	24 (86)	24 (69)
CQ	10 (16)	2 (7)	8 (22)
Both	5 (8)	2 (7)	3 (8)
Planned sample size*, median (IQR)	360 (150 to 672)	500 (218 to 1350)	267 (120 to 483)
Trial status, n (%)			
Completed	12 (19)	11 (39)	1 (3)
Discontinued	15 (24)	7 (25)	8 (23)
Not yet recruiting	7 (11)	0	7 (20)
Recruiting	29 (46)	10 (36)	19 (54)
Location, n (%)			
Africa	3 (5)	1 (4)	2 (6)
Asia	23 (37)	8 (29)	15 (43)
Europe	17 (27)	8 (29)	9 (26)
International	6 (10)	4 (14)	2 (6)
North America	10 (16)	4 (14)	6 (17)
South America	4 (6)	3 (11)	1 (3)
Placebo control, n (%)	30 (48)	11 (39)	19 (54)
More than two arms, n (%)	27 (44)	10 (37)	17 (49)
Patient setting, n (%)			
ICU	1 (2)	1 (4)	0
Inpatient	46 (73)	22 (79)	24 (69)
Outpatient	12 (19)	5 (18)	7 (20)
Unclear	4 (6)	0	4 (11)
Blinding, n (%)			
None	32 (51)	15 (54)	17 (49)
Outcome Assessor	1 (2)	1 (4)	0
Participant	3 (5)	1 (4)	2 (6)
Participant, Caregiver	26 (41)	11 (39)	15 (43)
Participant, Outcome Assessor	1 (2)	0	1 (3)

Abbreviations: chloroquine (CQ), hydroxychloroquine (HCQ), intensive care unit (ICU), interquartile range (IQR)

* Data were extracted from trial registries or publications. ** Including centers in multiple countries.

Supplements to “Mortality outcomes with hydroxychloroquine and chloroquine in COVID-19: an international collaborative meta-analysis of randomized trials”

- 1. Search strategy**
- 2. Template for invitation email**
- 3. Table S1. Group-level characteristics of unavailable trials**
- 4. Table S2A. Subgroup analyses**
- 5. Table S2B. Sensitivity analyses**
- 6. Figures S1A-C. Forest plots for sensitivity analyses**

Supplement 1. Search strategy

The COVID-evidence database includes trials registered on ClinicalTrials.gov or the WHO International Clinical Trials Registry Platform up to June 11, 2020, as well as trials published on the following sources up to April 9, 2020: PubMed, medRxiv, biorXiv, the WHO COVID-19 literature database, and a listing of all trials with ethical approval in Switzerland (for details please see the COVID-evidence study protocol on the Open Science Framework: <http://dx.doi.org/10.17605/OSF.IO/GEHFX>).

This Supplement describes the search strategy used to complement the COVID-evidence database with trials registered or published after April 9, 2020.

PubMed and the Cochrane COVID-19 trial registry were searched from inception to June 11, 2020. Search terms included extensive controlled vocabulary and Medical Subject Headings (MeSH). Search terms were the following: corona[ti] OR covid*[ti] OR sars[ti] OR severe acute respiratory syndrome[ti] OR ncov*[ti] OR "severe acute respiratory syndrome coronavirus 2" [Supplementary Concept] OR "COVID-19" [Supplementary Concept] OR (wuhan[tiab] AND coronavirus[tiab]) OR (wuhan[tiab] AND pneumonia virus[tiab]) OR COVID19[tiab] OR COVID-19[tiab] OR coronavirus 2019[tiab] OR SARS-CoV-2[tiab] OR SARS2[tiab] OR SARS-2[tiab] OR "severe acute respiratory syndrome 2"[tiab] OR 2019-nCoV[tiab] OR (novel coronavirus[tiab] AND 2019[tiab]) NOT (animals[mesh] NOT humans[mesh]) AND ("2019/12/01"[EDAT] : "3000/12/31"[EDAT])
AND
(((hydroxychloroquine[MeSH Terms]) OR (chloroquine[MeSH Terms])) OR (hydroxychloroquine[Title/Abstract])) OR (chloroquine[Title/Abstract])
AND
(randomized controlled trial[pt] OR controlled clinical trial[pt] OR randomized[tiab] OR placebo[tiab] OR clinical trials as topic[mesh:noexp] OR randomly[tiab] OR trial[ti] NOT (animals[mh] NOT humans [mh]))

The search was updated on October 16, 2020.

Supplement 2. Template for invitation email

Subject: Invitation to co-author a large-scale international collaborative meta-analysis on mortality in COVID-19 trials

Dear Dr. <last name>,

We are currently conducting a large-scale international collaborative meta-analysis on mortality in all ongoing or completed randomized clinical trials evaluating hydroxychloroquine or chloroquine for COVID-19. We are inviting all research groups worldwide testing these drugs to provide urgently needed evidence. We have no commercial interest with this work and aim to rapidly publish the results in a peer-reviewed journal. **Your core team is invited to co-author the publication.**

We use our COVID-evidence database (www.covid-evidence.org) for this work. COVID-evidence is supported by the Swiss National Science Foundation (Project ID 196190) and a large collaboration of researchers from Switzerland, the US, China, Canada, UK, France, Germany, Austria, Sweden, Netherlands, and other countries.

Our registered protocol can be found **attached** as well as registered on the Open Science Framework: [[link](#)]. Trials that are eligible for this project can be found at the end of the protocol.

Your study <url> is of high importance for this project. We would like to ask you how many patients died in your trial (see short questions below). We will use standard methods of meta-analysis and focus only on group-level (aggregated) mortality (no individual patient data needed). We will not do an in-depth review of the included trials as we aim to rapidly provide results, ideally from all trials worldwide. We describe the details of the study in the attached protocol.

For the meta-analysis, we kindly ask you to answer the following questions before July 7. If you are interested in collaborating, please let us know of your interest as soon as possible.

Question 1: Could you please confirm that these criteria apply to your trial?

- The trial is randomized and started enrollment before June 1, 2020
- The trial has at least one group of patients who receive hydroxychloroquine or chloroquine
- The trial has at least one control group that does not receive hydroxychloroquine or chloroquine

Question 2: For each of your study arms,

- What intervention did this group receive?
- How many patients were randomized to this group?
- Of these patients, how many have died?
- Of these patients, for how many it is unknown if they are dead or alive?

Please note that we are interested in these raw numbers regardless of the results of any statistical test. The numbers will be used to finalize the manuscript described in our registered protocol (attached).

We strive for a rapidly available, maximally informative publication with full transparency on methods. **With this publication we aim to make sure that all clinical trial data hitherto collected (unpublished or published) will be of use, regardless of whether the target sample size of each trial was reached or not. We invite your core investigator team as co-authors.** The manuscript will be shared with all co-authors for comments and the finalized manuscript will be uploaded as a preprint at medRxiv in parallel with submission to a peer-reviewed medical journal such as JAMA or the BMJ.

Thank you for considering our request! We kindly ask for your answer **before July 7**. If you are interested in collaborating, but are uncertain whether the data may be shared before July 7, please respond as soon as possible. Should you have any questions or comments, please let us know.

Best regards,

Cathrine Axfors, Andreas Schmitt, David Moher, Steve Goodman, John Ioannidis and Lars Hemkens for the COVID-evidence team

www.covid-evidence.org

Table S1. Group-level characteristics of randomized clinical trials evaluating hydroxychloroquine or chloroquine as treatment for COVID-19 not included in the meta-analysis.

Register ID	Contact name	Arms (n)	Treatment comparisons		Targeted sample size	Location	Setting	Trial status
			Experimental group (HCQ or CQ)	Control group				
NCT04315948	Ader	4	HCQ	SoC	3100	International	Inpatient	Recruiting
NCT04391127	Arreola Guerra	5	HCQ	Placebo	200	Mexico	Inpatient	Recruiting
NCT04351516	Bitzer	2	HCQ	Placebo	350	Germany	Outpatient	Recruiting
NCT04359953	Blanc	4	HCQ	Soc	1600	France	Inpatient	Not yet recruiting
ISRCTN86534580	Butler	2	HCQ	SoC	3000	Europe	Inpatient	Recruiting
ChiCTR2000029939	Cai	2	CQ	SoC	100	China	Inpatient	Recruiting
EUCTR2020-001271-33	CHU Angers study team	2	HCQ	Placebo	1300	France	Inpatient	Completed
EUCTR2020-001270-29	Sanofi Aventis study team	2	HCQ	SoC	350	International	Inpatient	Discontinued
ChiCTR2000030031	Deng	2	CQ	Placebo	120	China	Inpatient	Discontinued
NCT04331600	Duda-Sikula	2	CQ	SoC	400	Poland	Inpatient	Recruiting
NCT04328272	Farooq	3	HCQ	Placebo	75	Asia	Unclear	Not yet recruiting
NCT04339816	František	3	HCQ	Placebo	240	Europe	Inpatient	Recruiting
NCT04353037	Griffin	4	HCQ	Placebo	850	United States	Outpatient	Recruiting
NCT04349592	Harris	3	HCQ + Placebo	Placebo	456	Qatar	Inpatient	Not yet recruiting
ChiCTR2000029761	Huang	4	HCQ (3 arms with 3 different dosis)	SoC	240	China	Inpatient	Discontinued
ChiCTR2000029762	Huang	2	HCQ	SoC	60	China	Inpatient	Discontinued
EUCTR 2020-001469-35	Jankowska	2	CQ + telemedicine	telemedicine	400	Poland	Outpatient	Recruiting
NCT04354428	Johnston	3	HCQ + Placebo	Placebo + Placebo	630	United States	Outpatient	Recruiting
NCT04307693	Kim	3	HCQ	SoC	150	Asia	Inpatient	Discontinued
NCT04394442	Lutfy	2	HCQ	SoC	200	Saudi Arabia	Unclear	Recruiting
ChiCTR2000029837	Mao	2	CQ	Placebo	120	China	Inpatient	Discontinued
ChiCTR2000029826	Mao	2	CQ	Placebo	45	China	Inpatient	Discontinued
EUCTR 2020-001587-29	Menéndez	2	HCQ	Placebo	714	Spain	Inpatient	Recruiting
NCT04329611	Metz	2	HCQ	Placebo	1660	Canada	Inpatient	Recruiting

NCT04332991	Oldmixon	2	HCQ	Placebo	510	United States	Inpatient	Discontinued
NCT04342169	Pacchia	2	HCQ	Placebo	400	United States	Outpatient	Recruiting
NCT04353271	Richards	2	HCQ	Placebo	58	United States	Outpatient	Not yet recruiting
NCT04351191	Sarwar	4	HCQ (2arms with different dosis) CQ	Placebo Placebo	400	Pakistan	Unclear	Recruiting
NCT04346667	Sarwar	4	HCQ (2 arms with different dosis) CQ	Placebo Placebo	400	Pakistan	Outpatient	Recruiting
NCT04330586	Song	3	HCQ + Ciclesonide	Ciclesonide	141	Korea	Inpatient	Not yet recruiting
ChiCTR2000030987	Tong	3	CQ + Favipiravir	Favipiravir	150	China	Inpatient	Recruiting
EUCTR 2020-001265-36	UCD study team	3	HCQ	SoC	267	Ireland	Unclear	Recruiting
ChiCTR2000030417	Xu	2	CQ	Placebo	30	China	Inpatient	Discontinued
ChiCTR2000029740	Zhang	2	HCQ	SoC	78	China	Inpatient	Recruiting
ChiCTR2000029992	Zhenyu	3	HCQ CQ	SoC SoC	100	China	Inpatient	Not yet recruiting

* Trial includes more treatment arms than reported here; target sample size refers to all arms. Abbreviations: Chloroquine (CQ), hydroxychloroquine (HCQ), standard of care (SoC). Trial marked as “International” involves centers in multiple countries.

Table S2A. Subgroup analyses for random-effects meta-analysis on mortality for treatment of COVID-19 with Hydroxychloroquine.

Subgroup	Trials (n)	Patients (n)	OR (95% CI)	P value, test for interaction
Setting				
ICU	1	142	1.04 (0.49, 2.18)	0.98
Inpatient	21	9062	1.11 (1.02, 1.21)	
Outpatient	4	808	1.01 (0.06, 16.28)	
Published				
Yes	14	8981	1.12 (1.08, 1.16)	0.23
No	12	1031	0.92 (0.63, 1.34)	
Control				
Standard of care	17	8911	1.12 (1.04, 1.21)	0.15
Placebo	9	1101	0.88 (0.55, 1.41)	
COVID-19 diagnostic confirmation				
Confirmed	22	4215	1.11 (0.91, 1.36)	0.94
Confirmed or suspected	4	5797	1.1 (1.07, 1.14)	
Dose*				
High	3	6711	1.12 (1.01, 1.25)	0.29
Low	23	3301	0.97 (0.73, 1.30)	

* High dose: ≥ 1600 mg on day 1 and ≥ 800 mg from day 2. Low dose: < 1600 mg on day 1 or < 800 from day 2.

Table S2B. Sensitivity analysis for meta-analysis for mortality for treatment of COVID-19 with Hydroxychloroquine using different methods of combination.

	Reciprocal of contrasting arm added to zero events	Excluding trials with zero deaths in one or both arms	Trials with more than 50 participants*	Arcsine difference
	n = 26	n = 12	n = 15	n = 26
	OR (95% CI), I², Tau²			ASD (95% CI), I², Tau²
HKSJ-PM	1.11 (1.02, 1.20); 0,0	1.10 (1.03, 1.18); 0,0	1.10 (1.02, 1.19); 0,0	0.01 (0, 0.03); 0,0
HKSJ-SJ	1.06 (0.80, 1.40); 0,0.44	1.08 (0.96, 1.21); 0,0.02	1.04 (0.88, 1.22); 0,0.09	0 (-0.02, 0.03); 0,0
MH-DSL	1.11 (0.98, 1.24); 0,0	1.10 (0.98, 1.24); 0,0	1.10 (0.98, 1.24); 0,0	0.01 (-0.01, 0.03); 0,0
Peto	1.11 (0.98, 1.25); 0,0	1.10 (0.98, 1.24); 0,0	1.10 (0.98, 1.24); 0,0	0.01 (-0.01, 0.03); 0,0

HKSJ-PM = Hartung-Knapp-Sidik-Jonkman adjustment for random effects model, Paule-Mandel estimator for tau²;
HKSJ-SJ = Hartung-Knapp-Sidik-Jonkman adjustment for random effects model, Sidik-Jonkman estimator for tau²;
MH-DSL = Mantel-Haenszel method for random effects model, DerSimonian-Laird estimator for tau²;
PETO = Peto method with random effects model, Paule-Mandel estimator for tau²;

* Zero events were corrected by adding the reciprocal of the size of the contrasting study arm

Figure S1A. Forest plot of HKSJ-SJ model. Zero events were corrected by adding the reciprocal of the size of the contrasting study arm.

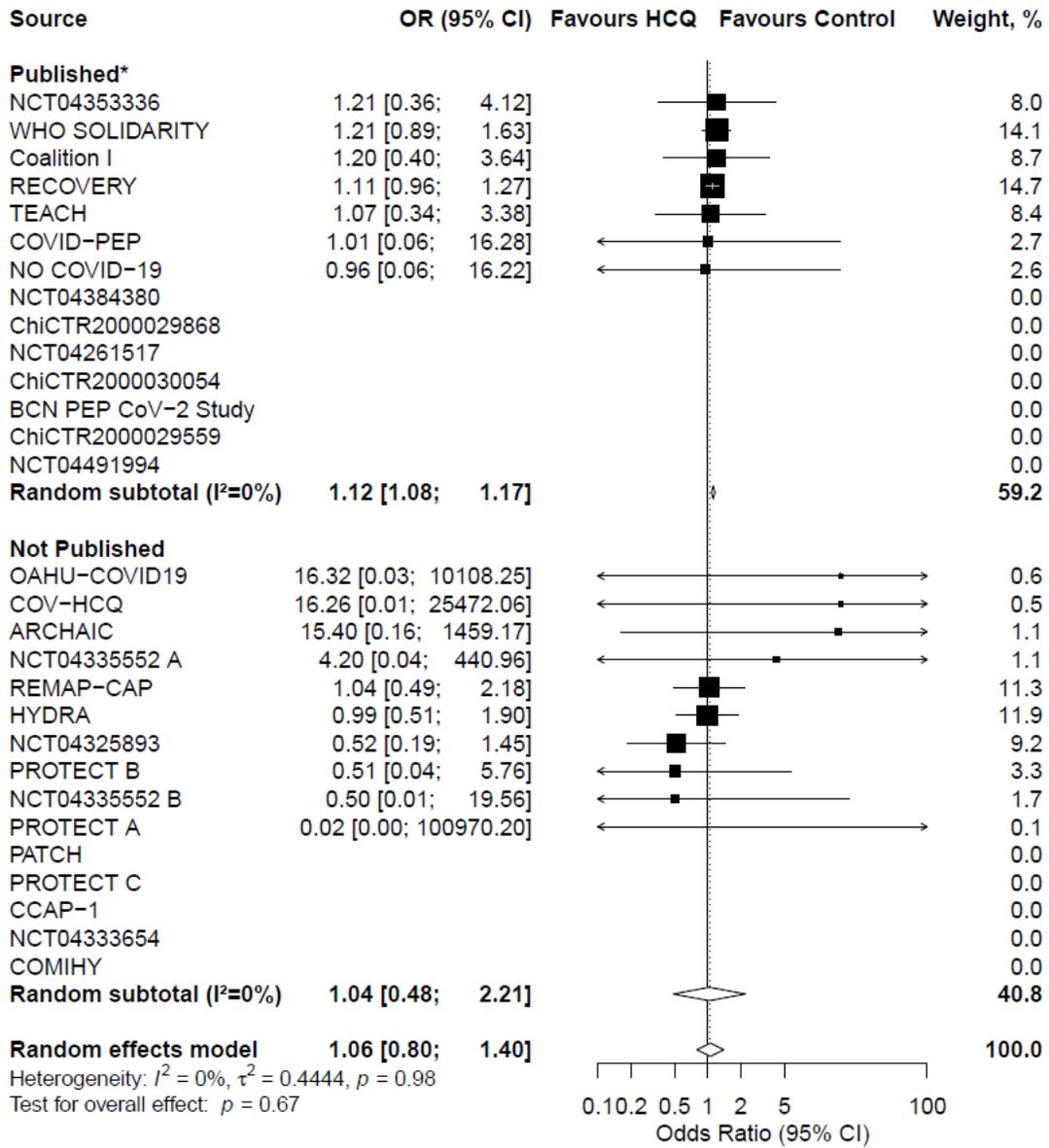


Figure S1B. Forest plot of MH-DSL model. Zero events were corrected by adding the reciprocal of the size of the contrasting study arm.

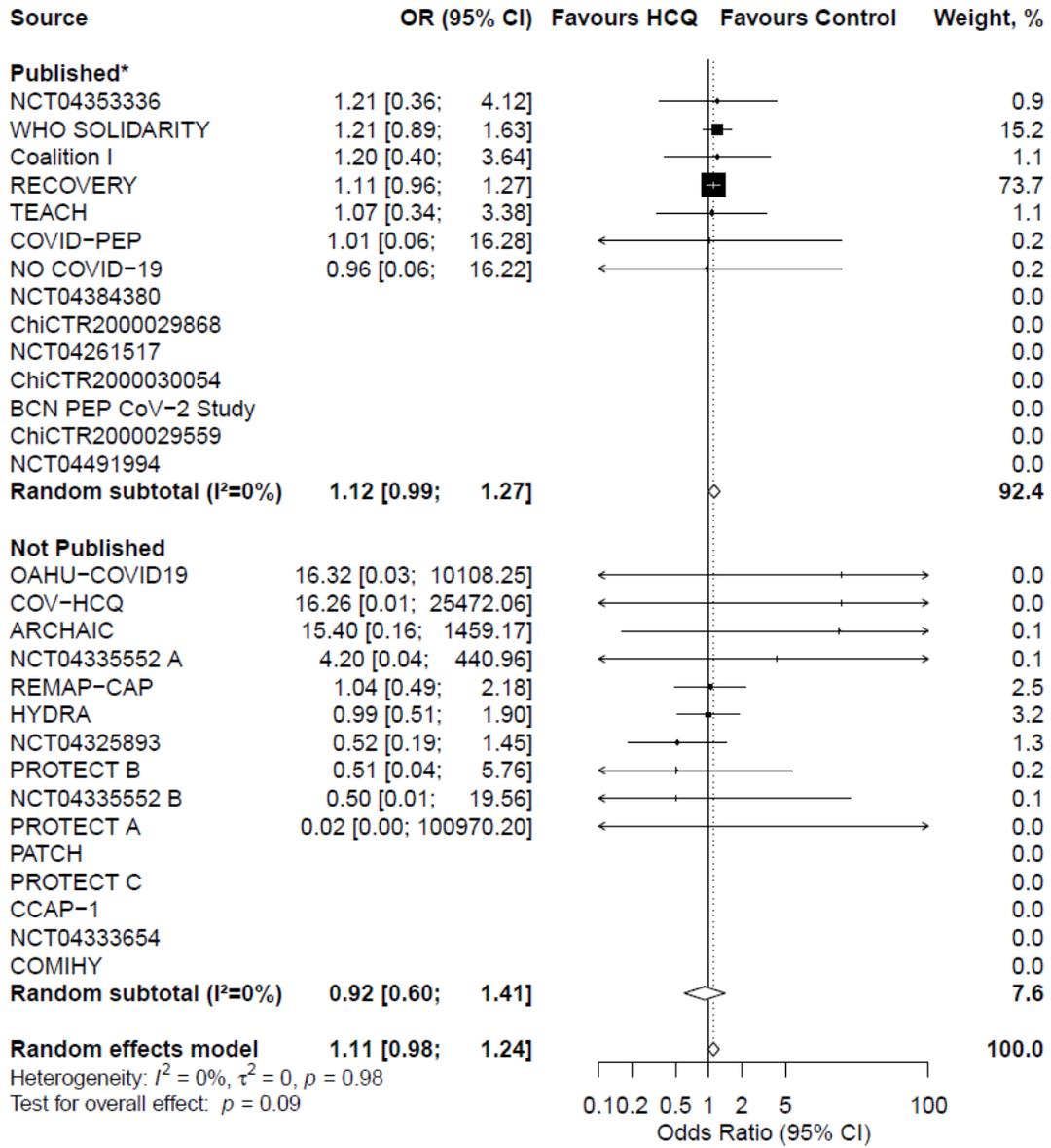


Figure S1C. Forest plot of Peto model. Zero events were corrected by adding the reciprocal of the size of the contrasting study arm.

