

## Supplementary Appendix

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## Data Safety and Monitoring Board

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Table S1. Risk Factors for Severe Disease

Medications and Biologic Therapies
Prednisone $\geq 7.5$ mg daily x 3 weeks (or equivalent) Methotrexate (Greater than or equal to 7.5 -15 mg weekly suggested) Azathioprine Cyclophosphamide within the previous 6 months Mitoxantrone Cell depleting therapy within the previous 24 months: cladribine Anti-TNF: infliximab, adalimumab, golimumab, etanercept, certolizumab Anti-IL17: secukinumab, ixekizumab, brodalumab mTOR inhibitors: sirolimus, everolimus Mycophenolate mofetil: mycophenolic acid Anti-IL12/23: Ustekinumab, risankizumab, guselkumab Anti-CD28: abatacept JAK2 inhibitors: tofacitinib, baricitinib, upadacitinib Anti-CD20: rituximab, ocrelizumab within the previous 12 months S1P inhibitors: fingolimod Anti-alpha4beta7: vedolozimab Anti-IL4: dupilumab Anti-IgE FcR: omalizumab
Medical Conditions and Other Risk Factors
Age 40 or over BMI >40 (calculated by self-report height and weight) Hypertension (on medical treatment) Current cigarette smoker Bone Marrow Transplant within previous 12 months Solid Organ Transplant AIDS/HIV CD4 <200 within last 6 months or CD4>200 but not on treatment Moderate Lymphopenia (within previous 6 months: Adults <500) Chronic Kidney Disease (eGFR < 60 including people on dialysis) Diabetes (on a hypoglycemic or insulin) Coronary Artery Disease (non-revascularized and as per physician diagnosis in medical chart) Heart Failure/Reduced LVEF (as per physician diagnosis in medical chart) Chronic Lung Disease (COPD, Asthma, interstitial lung disease, as per physician diagnosis) Any Current Cancer diagnosis (as per physician diagnosis) Acquired or Congenital Immune Deficiency (as per physician diagnosis in medical chart) Cirrhosis (normal INR and bilirubin and no history of ascites, encephalopathy, or variceal bleeding as per history and medical chart) Homelessness

Table S2. Modified Tisdale Scale

<b>Age ≥68 y</b>	<b>1</b>
<b>Female sex</b>	<b>1</b>
<b>Loop diuretic use: furosemide (Lasix), bumetanide (Bumex), torsemide</b>	<b>1</b>
<b>Serum K<sup>+</sup> ≤3.5 mEq/L on any blood test in the last 30 days</b>	<b>2</b>
<b>Admission QTc ≥450 ms on any recent ECG in the last 1 year</b>	<b>2</b>
<b>Acute MI</b>	<b>3</b>
<b>≥2 QTc-prolonging drugs (Including hydroxychloroquine)</b>	<b>3</b>
<b>Sepsis</b>	<b>3</b>
<b>Heart failure</b>	<b>3</b>
<b>One QTc-prolonging drug (This will be hydroxychloroquine)</b>	<b>3</b>

Modified to be suitable for outpatients who were screened using history and historical medical records. Modifications: (1) a low serum potassium within the previous 30 days (if any) replaced an admission potassium, and (2) an ECG within the previous year (if any) replaced an admission ECG.

Table S3. Use of telephone translation services during the trial

<b>Language</b>	<b>Number of Calls<sup>¶</sup></b>	<b>Language</b>	<b>Number of Calls<sup>¶</sup></b>
Tigrinya	36	French*	2
Tagalog*	27	Dinka	1
Spanish*	25	Nepali	1
Somali	19	Fuzhou	1
Punjabi*	18	Swahili	1
Vietnamese	17	Ukrainian	1
Mandarin	11	Uzbek	1
Cantonese	9	Malayalam	1
Arabic	8	Polish	1
Cambodian	7	Russian	1
Amharic	7	German*	0
Oromo	7	Czech*	0
Hindi*	5	Croatian*	0
Farsi*	4	Bosnian*	0
Portuguese*	4	Serbian*	0
Urdu*	3	Marathi*	0
Burmese	2	<i>Total Calls</i>	<b>220</b>

\*A researcher was also available to speak this language.

¶Participants often required more than one call throughout the trial.

Table S4. Primary and secondary outcomes in the per-protocol population (N=105)

<b>Outcome</b>	<b>Hydroxychloroquine (N=74)</b>	<b>Placebo (N=31)</b>	<b>P-value</b>
<b>Primary Outcome<sup>1</sup> - n (%)</b>	1 (1.4)	0 (0.0)	1.00*
<b>Secondary Outcomes</b>			
<b>Days to COVID-19 recovery - median (95% CI)<sup>2</sup></b>	12 (9-23)	13 (8-24)	0.56 <sup>†</sup>
<b>Disposition at 30 days - n (%)</b>			NC
<b>Recovered</b>	52 (70.3)	25 (80.6)	
<b>Ongoing symptoms, not hospitalized</b>	18 (24.3)	6 (19.4)	
<b>Unknown, not hospitalized or deceased</b>	4 (5.4)	0 (0.0)	
<b>Mortality within 30 days - n (%)</b>	0 (0.0)	0 (0.0)	NC
<b>Admission to ICU within 30 days - n (%)</b>	1 (1.4)	0 (0.0)	NC
<b>Hospitalization within 30 days - n (%)</b>	1 (1.4)	0 (0.0)	NC

\*Two-sided Fischer's exact test; <sup>†</sup>Log rank test.

<sup>1</sup>Hospitalization, invasive mechanical ventilation, or death within 30 days of treatment initiation.

<sup>2</sup>Excludes 8 subjects with missing symptom duration data in the hydroxychloroquine group

NC: secondary outcomes were not compared between groups following the pre-specified protected hierarchy.