

MATH+ HOSPITAL TREATMENT PROTOCOL FOR COVID-19

MEDICATION	INDICATION/INITIATION	RECOMMENDED DOSING	TITRATION/DURATION
Methylprednisolone	A. <i>Mild hypoxemia: requires O₂ via NC to maintain saturation > 92%</i>	40 mg IV bolus then 20 mg IV twice daily	A1. Once off O ₂ , then taper with 20 mg daily × 3 days then 10 mg daily × 3 days, monitor CRP response. A2. If FiO ₂ , or CRP increase move to B.
	B. <i>Moderate–severe hypoxemia (High Flow O₂, NIPPV, IMV)</i>	COVID-19 Respiratory Failure protocol (see flccc.net/respiratory-support-c19/) Preferred: 80 mg IV bolus, followed by 80 mg / 240 ml normal saline IV infusion at 10 ml/hr Alternate: 40 mg IV twice daily	B1. Once off IMV, NPPV, or High flow O ₂ , decrease to 20 mg twice daily. Once off O ₂ , then taper with 20 mg/day × 3 days then 10 mg/day × 3 days. B2. If no improvement in oxygenation in 2–4 days, double dose to 160 mg/daily. B3. If no improvement and increase in CRP/Ferritin, move to “Pulse Dose” below.
	C. <i>Refractory Illness/ Cytokine Storm</i>	“Pulse” dose with 125 mg IV every 6–8 hours	Continue × 3 days then decrease to 80 mg IV/daily dose above (B). If still no response or CRP/Ferritin high/rising, consider “Salvage Therapy” below
Ascorbic Acid	O ₂ < 4 L on hospital ward	500–1000 mg oral every 6 hours	Until discharge
	O ₂ > 4 L or in ICU	1.5–3 g intravenously every 6 hours	Sooner of 7 days or discharge from ICU, then switch to oral dose above
Thiamine	ICU patients	200 mg IV twice daily	Sooner of 7 days or discharge from ICU
Heparin (LMWH)	Hospital ward patients on ≤ 4 L O ₂	0.5 mg/kg twice daily Monitor anti-Xa, target 0.2–0.5 IU/ml	Until discharge then start DOAC at half dose × 4 weeks
	ICU patients or > 4 L O ₂	1 mg/kg twice daily Monitor anti-Xa levels, target 0.6–1.1 IU/ml	Later of: discharge from ICU or off oxygen, then decrease to hospital ward dosing above
Ivermectin (should be considered a core medication)	Upon admission to hospital and/or ICU	0.3 mg/kg per dose – one dose daily, minimum of 2 days	continue daily until recovered (max 5 days)
Vitamin D	Hospital ward patients on ≤ 4 L O ₂	Calcifediol preferred: 0.532 mg PO day 1, then 0.266 mg PO day 3 and 7 and weekly thereafter Cholecalciferol: 10,000 IU/day PO or 60,000 IU day 1, 30,000 IU days 3 and 7 and then weekly	Until discharge
	ICU patients or on > 4 L O ₂	Cholecalciferol 480,000 IU (30 ml) PO on admission, then check Vitamin D level on day 5, if < 20 ng/ml, 90,000 PO IU/day × 5 days	Until discharge from ICU
Atorvastatin	ICU Patients	80 mg PO daily	Until discharge
Melatonin	Hospitalized patients	6–12 mg PO at night	Until discharge
Zinc	Hospitalized patients	75–100 mg PO daily	Until discharge
Famotidine	Hospitalized Patients	40–80 PO mg twice daily	Until discharge
Therapeutic Plasma Exchange	Patients refractory to pulse dose steroids	5 sessions, every other day	Completion of 5 exchanges

Legend: CRP = C-Reactive Protein, DOAC = direct oral anti-coagulant, FiO₂ = Fraction of inspired oxygen, ICU = Intensive Care Unit, IMV = Invasive Mechanical Ventilation, IU = International units, IV = Intravenous, NIPPV = Non-Invasive Positive Pressure Ventilation, O₂ = oxygen, PO (per os) = oral administration

MATH+ HOSPITAL TREATMENT PROTOCOL FOR COVID-19

TO CONTROL INFLAMMATION & EXCESS CLOTTING

In all COVID-19 hospitalized patients, the therapeutic focus must be placed on early intervention utilizing powerful, evidence-based therapies to counteract:

- The overwhelming and damaging inflammatory response
- The systemic and severe hyper-coagulable state causing organ damage

By initiating the protocol soon after a patient meets criteria for oxygen supplementation, the need for mechanical ventilators and ICU beds will decrease dramatically.

TREATMENT OF LOW OXYGEN

- If patient has low oxygen saturation on nasal cannula, initiate heated high flow nasal cannula.
- Do not hesitate to increase flow limits as needed.
- Avoid early intubation that is based solely on oxygen requirements. Allow “permissive hypoxemia” as tolerated.
- Intubate only if patient demonstrates excessive work of breathing.
- Utilize “prone positioning” to help improve oxygen saturation.

ABOUT THE MATH+ HOSPITAL TREATMENT PROTOCOL FOR COVID-19

Our **MATH+** protocol is designed for hospitalized patients, to counter the body’s overwhelming inflammatory response to the SARS-CoV-2 virus. The protocol is based on numerous medical journal publications over decades. It is the hyper-inflammation, not the virus itself, that damages the lungs and other organs and ultimately causes death in COVID-19. We have found the **MATH+** protocol to be a highly effective combination therapy in controlling this extreme inflammatory response and we have now added ivermectin as a core component given the profound emerging efficacy data in hospitalized patients reviewed here (www.flccc.net/flccc-ivermectin-review-covid-19).

The steroid Methylprednisolone is a key component, increasing numbers of studies (see <https://flccc.net/medical-evidence>) show its profound effectiveness in COVID-19, which is made more potent when administered intravenously with high doses of the antioxidant Ascorbic acid given that the two medicines have multiple synergistic physiologic effects. Thiamine is given to optimize cellular oxygen utilization and energy consumption, protecting the heart, brain, and immune system. The

anticoagulant Heparin is important for preventing and dissolving blood clots that appear with a very high frequency in patients not given blood thinners. The **+** sign indicates several important co-interventions that have strong physiologic rationale and an excellent safety profile. It also indicates that we plan to adapt the protocol as our insights and the published medical evidence evolve.

Timing is a critical factor in the successful treatment of COVID-19. Patients must go to the hospital as soon as they experience difficulty breathing or have a low oxygen level. The **MATH+** protocol then should be administered soon after a patient meets criteria for oxygen supplementation (within the first hours after arrival in the hospital), in order to achieve maximal efficacy as delayed therapy has led to complications such as the need for mechanical ventilation.

If administered early, this formula of FDA-approved, safe, inexpensive, and readily available drugs can eliminate the need for ICU beds and mechanical ventilators and return patients to health.

DISCLAIMER

This protocol is solely for educational purposes regarding potentially beneficial therapies for COVID-19. Never disregard professional medical advice because of something you have read on our website and releases. It is not intended to be a substitute for professional medical advice, diagnosis, or treatment in regards to any patient. Treatment for an individual patient should rely on the judgement of your physician or other qualified health provider. Always seek their advice with any questions you may have regarding your health or medical condition.

CONTACT

FLCCC Alliance
6006 N Highlands Avenue
Madison, WI 53705-0000

Physician Contact
Howard Kornfeld, MD — support@flccc.net
Keith Berkowitz, MD — keith@centerforbalancedhealth.com

Media Relations
press@flccc.net



For updates, references and more information please see

www.flccc.net



TO CONTROL INFLAMMATION & EXCESS CLOTTING

In all COVID-19 hospitalized patients, the therapeutic focus must be placed on early intervention utilizing powerful, evidence-based therapies to counteract:

- The overwhelming and damaging inflammatory response
- The systemic and severe hyper-coagulable state causing organ damage

By initiating the protocol soon after a patient meets criteria for oxygen supplementation, the need for mechanical ventilators and ICU beds will decrease dramatically.

MATH+ PROTOCOL

[Only for use in hospitals in the treatment of COVID-19]

1. **Methylprednisolone** [Intravenous]

- A. Mild hypoxia (< 4L): 40 mg daily until off oxygen
 - B. Moderate-severe illness: 80 mg bolus, then 20mg q6h IV push for 7 days*
 - Alternate: 40 mg q12h for 7 days*
 - Day 8: Switch to oral prednisone, taper over 6 days
- *Consider higher doses for patients with non-improving ARDS/oxygenation and/or with persistent, rising, or severely elevated inflammatory markers (cytokine storm), i.e. 60-125 mg q6h-q8h, or 1,000mg/day for 3 days

2. **Ascorbic Acid** [High Dose Infusion]

- 3 grams / 100 ml – q6h
- Continue for a total of 7 days or until discharged

q6h/q8h/q12h = every 6/8/12 hours
1 mg Heparin = 500 int. units (IU)
CrCl = Creatinine Clearance (Cr)

3. **Thiamine**

- 200 mg IV – q12h – until discharged

4. **Heparin** [Low Molecular Weight Heparin / LMWH]

- A. Stable patient on medical floor/ward: 0.5mg/kg q12h; if CrCl ≤ 30 ml/min, give once a day
- B. Critically ill or ICU patient: 1mg/kg q12h unless contraindicated, dose adjust for CrCl 15-30 ml/min
 - If CrCl ≤ 15 ml/min, use unfractionated heparin [UFH]
 - Monitor antifactor-Xa activity, target range is 0.6-1.1 units/ml
- Continue until discharged

5. **PLUS** optional co-interventions: Melatonin (6-12 mg at night), Zinc (75-100mg/day), Vitamin D3

TREATMENT OF LOW OXYGEN

- If patient has low oxygen saturation on nasal cannula, initiate heated high flow nasal cannula.
- Do not hesitate to increase flow limits as needed.
- Avoid early intubation that is based solely on oxygen requirements. Allow “permissive hypoxemia” as tolerated.
- Intubate only if patient demonstrates excessive work of breathing.
- Utilize “prone positioning” to help improve oxygen saturation.

For updates, references and more information please see

www.flccc.net