sample quality, etc. In the setting of a negative test in high-suspicion opatient in isolation & repeat testing in 24h. PCR-NEGATIVE BUT HIGH COVID SUSPICION: A single FDA Variation is due to timing of test during course of illness, si alternative diagnosis) it is *strongly* recommended to keep

are

disease

mild-moderate COVID-19

rare (~3%). Antibiotics in

with COVID-19 are

bacterial co-infections

Early in disease,

UCDH ADULT COVID-19 TREATMENT GUIDELINES

DISEASE SEVERITY

INFECTIOUS DISEASE RECOMMENDATIONS

Ver. 15 7/29/21

- Recommend against the use of Dexamethasone.
- If all below criteria are met, consider EUA monoclonal antibodies (through outpatient monoclonal antibody therapy workflows)
 - No indication for admission
 - Mild-moderate disease w/ <10 days of influenza-like illness symptoms (no O2 requirement)
 - 1+ of the following risk factors: Age > 64, immunocompromised, COPD, HTN/CVD, CKD, DM, BMI > 34, pregnancy

- Prioritize clinical trial enrollment. Contact ID Clinical Trial coordinator to see if eligible (see COVID Clinical Trials criteria)
- Recommend against the use of Dexamethasone.
- If symptom onset < 10-14 days AND high risk for progression to severe disease (multiple risk factors for severe disease +/- CXR findings), start **Remdesivir** x 5 days at dose and duration discussed below.

Mechanical Ventilation, or ECMO)

- Prioritize clinical trial enrollment. Contact ID Clinical Trial coordinator to see if eligible (see COVID Clinical Trials criteria)
- Start Remdesivir 200mg IV for one day followed by Remdesivir 100mg IV once daily for 4 days or until hospital discharge, whichever comes first. Extending duration to 10 days in the setting of no clinical improvement should be done in consultation with ID. Most benefit if started within 10-14 days of symptom onset. Consider risks/benefits of starting Remdesivir if creatinine clearance < 30 mL/min (can discuss with ID or ID pharmacy). Not contraindicated in RRT.
- If progressive hypoxia or ≥ 3L/min NC, add Dexamethasone 6mg IV or PO daily for up to 10 days or until hospital discharge, whichever comes first.
- May consider adding Dexamethasone if stable disease on ≤ 2L/min AND patient has multiple risks for severe disease.
- Baricitinib may be considered as an alternative to Dexamethasone if there are contraindications to steroid use (i.e. DKA)

HFNC or Non-Invasive Ventilation

- Prioritize clinical trial enrollment. Contact ID Clinical Trial coordinator to see if eligible (see COVID Clinical Trials criteria)
- Use one of the following options:
- Dexamethasone plus Remdesivir at the doses and durations discussed above (if < 14 days of symptom onset)
- Dexamethasone alone at dose and duration discussed above (e.g., when combination therapy with Remdesivir cannot be used)
- If admitted within 24 hours (or within 48 hours w/ rapid progression) add, in combination w/ Dexamethasone, give Tociluzimab x 1
- •8mg/kg IV one time by IBW (max dose 800mg)
- Exclusion criteria: immunomodulating drugs, AST or ALT > 5x ULN, ANC < 500, plts < 50,000, hi risk of GI perf, hi suspicion bacterial or fungal infection

Hospitalized

Mechanical Ventilation or ECMO

- Start Dexamethasone at the dose and duration discussed above
- If admitted within 24 hours and no exclusion criteria met add, in combination with Dexamethasone, Tocilizumab at the dose discussed above
- No evidence to support using Remdesivir in this population; however, if ≤7 days of symptoms OR patient had been started on an alternate form of supplemental oxygen within the past 24h (NC, HFNC, or NIV) could consider addition of Remdesivir in consultation with ID.

Immunocompromised

• For the severely immunocompromised (heme-malignancy, SCT, SOT, HIV/AIDS, etc) consider ID consultation for additional guidance