### **ICU Management of COVID-19**

Desiree Wood, DNP, ARNP

VA Puget Sound Health Care System

## **Objectives**

- Clinical syndromes and severity of COVID-19
- When to admit to the ICU
- Initial evaluation
- Respiratory failure and intubation
- ARDS management
- Treatment standards

### Put on Your Own Mask First: COVID unknown





# The Virus vs The Disease

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## Clinical presentation of critically ill COVID-19

- Characteristic Lancet Resp Med (n=52) JAMA (n=24) NEJM (n=25)
- Demographics
   Age (mean, SD or range) 60 (SD 13) 70 (43-92) 64 (23-97) Male 67% 52% 63%
  Symptoms
  Fever 98% 52% 50%

- Cough 77% 48% 88%
  Dyspnea 64% 76% 88%
- Comorbidities
   Chronic cardiac disease 10% 43% -
- · COPD 8% 33% 4%
- Cerebrovascular disease 14% Diabetes 17% 33% 58%
- OBESE

Clinical presentation of critically ill COVID-19

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- "influenza like illness" fever, fatigue, cough anorexia malise, muscle pain sore throat dyspnea, nasal congestion or HA
- \*\*decrease taste, GI upset, diarrhea, ASYMPTOMATIC?!?! Pneumonia
- Adult with pneumonia on CXR, but no signs of severe pneumonia listed below Severe COVID/Pneumonia
- Suspected respiratory infection, plus elevated respiratory rate >30 with respiratory distress; or Sp02<94% at rest on ambient air; Pa02/Fi02 ≤300mmHg

WHO Clinical Management of SARI when COVID-19 is Suspected, March 13, 2020

### Clinical Syndromes and severity of COVID-19

#### ARDS

Bilateral opacities on CXR not fully explained by volume overload lobar lung collapse or nodules

-not clinically fully explained by cardiac failure or overload -PaO2: FiO2 ration <300 w/PEEP or CPAP >5

-Mild ARDS : P:F ratio =200-300

-Moderate ARDS: P:F ratio = 100-200 -Severe ARDS: P:F ratio <100

If Pa02 not available, Sp02/Fi02 <315 suggests ARDS

JAMA 2012;307:2526-253



When to admit?

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### Hospital vs ICU admission

 $\label{eq:Admit} \textbf{Admit patients with } \underbrace{\textbf{evidence of pneumonia, oxygen requirement}}$ 

- ICU admission
- Typical ICU admission criteria (respiratory failure, shock, etc.) Severe COVID pneumonia patients
- Evidence of cardiac disease due to COVID
- Uncertainties

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- Should certain patients with higher risk features for poor outcomes be admitted if they present with mild disease only?
- How long should patients with non-severe pneumonia remain in the hospital before discharging home out of concern for worsening disease?

### **Evaluation**

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Vital Signs: temp, HR, RR, BP, SpO2, Glasgow Coma Scale (confusion)

- Early warning severity score calculated by SOFA at 24 and 48 hours
- (respiratory variables, MAP, GCS, bilirubin, platelets, creatinine)
- COVID Extended Review of Systems: Fever (present in 99%); dyspnea;
- requirement; dry cough; rhinorrhea; myalgias; fatigue; diarrhea (can precede resp sx); nausea/vomiting; contacts with sick individuals in past 3 week

### Labs in COVID-19

- Laboratory studies: All patients should have COVID testing according to updated
- CDC or VHA guidance as test availability evolves
- • For patients with moderate or severe illness:
- $\bullet$  CBC w/differential, comprehensive metabolic panel (electrolytes, liver
- $\bullet \ \ function \ tests), \ magnesium, \ PT/PTT/INR$
- - Troponin, CRP, BNP; D-dimer (DIC panel); ferritin; LDH
- INFLAMATORY MARKERS
- - Chest X-ray; EKG
- $\bullet$   $\,\bullet$  For patients with severe illness, notify ICU team immediately

### **MICROBIOLOGY**

- Initial microbiologic evaluation and antimicrobials
- Nasopharyngeal swab more sensitive than oropharyngeal
- •Lower respiratory tract sample if feasible: but many labs don't have testing ability for this
- Do not induce sputum
- - Avoid bronchoscopy: aeroslized
- - Endotracheal aspirate for intubated patient (keep circuit closed)
- • Initial empiric antibiotics per CAP Guidelines
- • Evaluate for other infections: influenza testing, consider respiratory
- virus panel, procalcitonin (if available), blood cultures

### **Complications**

Complications Lancet Resp Med (n=52) JAMA (n=21)

- ARDS 67% to 95%
- Mechanical ventilation 64% to 71%
- Acute kidney injury 29% to 19%
  Renal replacement therapy 17% Not reported
- Cardiac injury 23% to 33%
- Liver dysfunction 29% to 14%
  Died (28-days) 62% at 28 days 52% (38% still in ICU at time of publication)

#### Clinically observed phases of COVID-19 from colleagues at Emory Critical Care Center

- Phase 1 "Viral prodrome" with non-specific symptoms, N/V, poor PO
- Phase 2 "Slow, smoldering hypoxia" pneumonia diagnosed at 4-5 days
- More comfortable but hypoxic, difficulty mobilizing thick secretions
- Phase 3 "The Struggle Bus"
- Increasing oxygen requirements, more cough, secretions, and worsening CXR
- Phase 4 "Respiratory Collapse"
- Intubation, relatively normal compliance, thick secretions
- Phase 5 "Rapid death death or steady resolution"
- · MOSF, hyper-inflammatory state

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### **Prognostication for poor outcomes?**

- Older age
- Comorbidities (hypertension, CVD, diabetes, COPD)
- Immune and inflammatory markers
- - Lower lymphocytes counts (particularly of CD4+ T-lymphocytes)
- - Higher leukocytes counts and neutrophil-lymphocyte-ratio (NLR)
- - Elevated inflammatory markers (IL6, IL1, TNF-alpha)
- -Immunosuppressed (transplant)

### Management in the ICU

- Bundle assessments to minimize personnel exposed, PPE used
- Think before ordering: avoid "routine" imaging studies and other tests
- Recommend placement of PICC line early in clinical course to facilitate IV access, lab draws
- After intubation, place arterial line given potential for rapid development of
- · hemodynamic instability.
- If patient does not have a PICC line, place central venous line at this time.

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### Non-invasive management of respiratory failure

- Recommend early intubation when requiring higher amounts of oxygen
- HFNC/NIPPV may not prevent intubation but rather delay it, with potential
- Initial NIPPV may in particular yield worse outcomes (unpublished data from Italy)
- Open systems may increase droplet dispersion (risk to HCW) with poorly fitting interface
- Proceed with early intubation if deteriorating respiratory, hemodynamic, or mental status to avoid emergent procedure
- Adequacy of resources (ventilators) may be a factor in some circumstances

### **Special considerations**

- · Patients who are DNI
- Keep flows for HFNC <30L/min; may trial increased FIO2/flow and consider transition to
- · comfort measures if failing for those who are DNI
- Ventilatory failure as predominant pathophysiology e.g. COPD exacerbation
- Use closed expiratory circuit mask/device with HEPA filter and ensure good mask seal
- • CPAP use by Helmet may further decrease aerosols (Not available at VA) • • Use airborne precautions if on HFNC or NIPPV
- Negative pressure room
- PAPR or N95s for staff

### When and how to intubate

- Early intubation e.g. once requiring >6L NC
- · Avoid bag mask ventilation
- Maximize pre-oxygenation with NC, simple FM, or non-rebreather
- RSI with video laryngoscopy by most experienced provider
- Preferred PPE: PAPR with shroud, gown, and gloves that extend over gown cuffs

### **ARDS management: LPV**

- Low tidal volume ventilation with 4-6 cc/kg of ideal body weight
- Up to 8 cc/kg allowed\* if needed for breath stacking, dysynchrony
- Remember, IBW is based on HEIGHT and sex, not actual weight Keep plateau pressures <30\*
- Target Spo2 92-96%
- Start with standards ARDSnet PEEP ladder
- Consider high PEEP strategy for more severe ARDS
- Particularly with low compliance, recruitable lung
- - Monitor hemodynamics MAP may drop (decreased venous return-> cardiac output

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### **ARDS Management continued**

- ng for moderate-severe ARDS for 12-16 hours a day recommended per SCCM
- guidelines for care of critically ill patient with COVID-19

  Severe ARDS (P:F<150 mm Hg), Fio2 of ≥ 0.6, PEEP of ≥ 5 cm of water, and TV of 6 cc/kg of IBW1
- Incorporate staff expertise, time, resources (use of PPE), and staff exposure in risk/benefit
- Reports of high compliance ARDS high PEEP and proning may not be as beneficial
- Possible role of inhaled pulmonary vasodilators (e,g, Flolan)
- Neuromuscular blockade not of benefit in general in ARDS
- Conservative fluid management and diuresis

### Monitoring for and treating cardiovascular complications

- Consider baseline TTE to document cardiac function given risk for
- COVID-19 cardiomyopathy : Especially in COVID-19 intubated patient
- Daily EKG, particularly if on hydroxychloroquine +/- azithromycin
- Monitor daily troponin, BNP

Troponin elevation is common, potentially due to:

- Myocardial injury or MI (type I plaque rupture vs. type II, stress)
- Stress cardiomyopathy Viral cardiomyopathy
- Increasing troponin and CK or decreasing ScVO2 could indicate development of cardiomyopathy/myocarditis
- Can appear late in the course, even after patients are recovering from ARDS
- May respond to dobutamine

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### Other systems affected

- Renal: AKI not related to any hemodynamic or volume status/early ATN
- Heme: Elevated D-Dimer consideration of anticoagulation (Italy)
- Neurologic: Marked encephalopathy with agitation and high sedation
- requirements has been observed (encephalitic component?) (Emory)
- Pulmonary recovery has preceded neurological recovery
- SAT to tolerate SBT is rate limiting step
- Avoid/minimize use of benzodiazepines: Propofol shortage!
- -skin, pressure ulcers, restraints, critical illness weakness

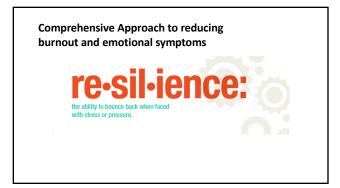
### Therapies ? This has changed every week @

Stage 1 "Early infection"

- - Supportive care
- Stage 2 "Pulmonary phase"
- · Stage 3 "Hyperinflammation phase"
- -NIH recommends against using hydroxychloroquine out side of clinical trials!
- -buzz around Remdesivir just FDA approved
- -steroids? IVIG? Again what is the evidence?

### Considerations

- · Limited family presence
- Code protocols
- Palliative care
- Rehabilitation
- Tracheostomy
   Is it fact or an opinion
- Goal directed therapies
- Stigma
- Testing
- Self care



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