COVID19: FOR THE ANESTHESIA/CRITICAL CARE PROVIDER

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The Anesthesia Provider's evolving Role

- Understand COVID19 Basic Science, Transmission, and PPE
- Treatment COVID19
 - Workup and Medicinal Therapies
 - Hypoxia and ARDS therapies
- Lots of information use slides as reference

GOALS AND OBJECTIVES



Disaster Planning OR to ICU Bed Conversion Protocol (Last Revision, March 19, 2020)

Locations: Filled according to the following priority levels 10 additional converted Operating Room to Intensive Care Unit beds (OR/ICU bed's) Priority 1 (i.e. filled first): OR 2, 3, 4, 5, 6, 7, 8, Priority 2: OLD OB OR 1 and 2 Priority 3: CV OR 7 Collectively known as the OR Block of ICU beds.

Physician Anesthesiologist and CRNA, Chain of Command

-Attending Intensivist will be responsible for admitting patients into the converted operating room/ ICU beds. He/she will be in charge of all aspects related to the care of the patient including appropriate oversight and delegation of various responsibilities to OR/ICU supervisors (see below for definition). -Resident Anesthesiologists report to their supervising physician under all circumstances -CRNA's when supervising OR/ICU rooms report directly to the attending intensivist. They will remain independent when performing regular anesthesia duties -Physician Anesthesiologist when supervising OR/ICU rooms will report to the intensivist. Anesthesiologist when supervising OR/ICU rooms will report to the intensivist. Anesthesiologist and anesthesiologist is the attending intensivist if appropriately cross trained. When the physician anesthesiologist is the attending intensivist, they must abdicate their role of OR/ICU room supervisor.

Staff Assignments

All staff members will report to the OR/ICU Room Supervisor or Attending Intensivist

1 ICU or CVICU charge nurse in charge of nursing for the OR Block of ICU beds

OR Nursing to flex to ICU Nurses 1 Float CCRN to 4 OR RN Model for consultative and break purposes, CCRN cannot be responsible for 2 priority locations, They cannot be assigned their own patients.

1:1 Patient to OR Nurse Ratio

- Minimize personal/ patient exposure to COVID 19 in the perioperative Setting
- OR's can be converted to Intensive Care Unit Beds (Italy, Washington State, New York, China etc.)
- ▶ We are all intensivist in the OR YOU CAN HELP
- Kaweah Delta is developing Protocols for Anesthesia to provide Intensive Care
- Anesthesia providers will need to adapt and become critical care providers

THE ANESTHESIA PROVIDER'S EVOLVING ROLE IN THIS PANDEMIC



Modified from the Ontario Health Plan for an Influenza Pandemic Workgroup. Critical Care During a Pandemic.

Source: SCCM Resource availability

https://sccm.org/Blog/March-2020/United-States-Resource-Availability-for-COVID-19?zs=jxpjd1&zl=w9pb6



SARS-COV2 VIRUS

- Causes the disease known as Covid19
- A Coronavirus Named after the characteristic solar flare like image seen on Electron Microscopy
- > SARS, MERS, and now COVID19
- ▶ positive sense RNA
- COVID19 primarily enters
 Angiotensin Converting Enzyme 2
 Receptor in lung epithelium
- Transmission via COVID-19 viruses in respiratory droplets

Image taken from Wikipedia: <u>https://en.wikipedia.org/wiki/Coronavirus</u>

CORONAVIRUS PANDEMIC

COVID-19 is an infectious disease caused by SARS-CoV-2, a new type of coronavirus detected in China in late 2019.



Data shows the disease is mild in 80 percent of patients, severe in 13 percent, and critical in 6 percent.

Most common symptoms:

Fever Fatigue Dry cough

Some patients may also have:

- Aches and pains
- Runny nose
- Sore throat
- Shortness of breath
- 🏓 Diarrhoea

In critical cases. COVID-19 can cause severe pneumonia or a multiple-organ failure and can lead to death.



Shenzhen-based family visit infected relatives in Wuhan, and return with illness







Transmission

$R_0 = 2-4$

COVID-19 is spread via **DROPLETS** and **FOMITES** in close spaces.

Airborne spread has not been reported but is theoretically possible when droplets become aerosolized.

Fecal shedding is possible but fecal-oral transmission is not believed to be a major route of transmission.

van Doremalen N, Bushmaker T, Morris DH, Holbrook MG, Gamble A, Williamson BN, et al. Aerosol and Surface Stability of SARS-CoV-2 as Compared with SARS-CoV-1. New England Journal of Medicine. 2020 Mar 17;0(0):null.

C Half-Life of Viable Virus



HALF LIFE OF VIRUS ON DIFFERENT SURFACES

> Airborne precautions: N95 mask and face protection

- Droplet precautions: surgical mask
- Contact precautions: gloves and gown, dedicated disposable equipment

PERSONAL PROTECTIVE EQUIPMENT

COVID-19 PPE GRID- What personal protective equipment to wear and when?

It is critical for all of us to apply critical thinking skills in using Standard Precautions, essentially donning PPE that is appropriate for situations in which we anticipate exposure to body fluids and respiratory secretions.

The CDC and Cal-OSHA/OSHA requires certain types of transmission based precautions for different scenarios. Based on these recommendations what follows is a table with appropriate precautions given a particular scenario...scenarios or questions not captured please contact command center **624-5000**.

Scenario	Transmission Based Precaution (Healthcare Personnel)	PPE required	Patient	Disinfect Surfaces	
Respiratory Symptom Patient in Waiting Room/Lobby/ED Triage this includes Clinics/Urgent Care.	Droplet/Contact	*Facemask; Yellow or White Isolation Gown; Gloves. PPE needed only if treating patient	Facemask	Purple top disinfectant (Offices only - QT3 disinfectant)	
Respiratory Symptom Patient in CT, MRI, Ultrasound, X-ray, nuclear medicine, Cath lab.	Droplet/Contact	*Facemask; Yellow or White Isolation Gown; Gloves	Facemask	Purple top disinfectant (Offices only - QT3 disinfectant)	
Respiratory Symptom patient suspected of COVID-19 Before specimen collection or after specimen collection completed.	Droplet/Contact	*Facemask; Yellow or White Isolation Gown; Gloves	Facemask	Purple top disinfectant (Offices only - QT3 disinfectant)	
Respiratory Symptom Patient with suspected or confirmed COVID-19 during specimen collection AND all other <u>Aerosol</u> <u>Generating Procedures</u> (during intubation, sputum induction, ET aspiration, bronchoscopy, nebulizer treatments, BiPap, HiFlo, extubation). Coordinate with Respiratory therapy for a hepa filter for use during the collection.	Airborne/Contact (HEPA filter or negative air pressure rooms)	N95 Mask; Yellow or White Isolation Gown; Gloves; Goggles or Face shield as needed	Facemask	Purple top disinfectant (Offices only - QT3 disinfectant)	
Confirmed positive COVID-19 patient	Droplet/Contact	Facemask; Yellow or White Isolation Gown; Gloves	N/A	Purple top disinfectant (Offices only - QT3 disinfectant)	
Cleaning COVID-19 patient room or specimen collection room	Droplet/Contact	**Facemask; Yellow or White Isolation Gown; Gloves	N/A	Purple top disinfectant (Offices only - QT3 disinfectant)	
Anesthesia (during intubation/LMA/extubation).	Airborne/Contact (Negative Pressure, HEPA filter, AIIR not needed)	N95 Mask; Yellow or White Isolation Gown; Gloves; Goggles or Face shield as needed	None	Oxivir disinfectant wipes	
Surgery/Endoscopy.	Droplet/Contact	*Facemask; Yellow or White Isolation Gown; Gloves	None	Oxivir disinfectant wipes	
Home Health/Private Home Care/Hospice/Community Outreach.	Droplet/Contact	*Facemask; Yellow or White Isolation Gown; Gloves	Facemask or None	Purple top disinfectant (Offices only - QT3 disinfectant)	

*PPE is required only when the patient is not wearing a Facemask or is unable/unwilling to contain his/her respiratory secretions. **PPE is only required if patient is present during room cleaning

Goggles or face shield as needed = use if secretions are expected White gown should be used when secretions or bodily fluid saturation is expected 3.19.20

Recommendations for Airway Management in a Patient with Suspected Coronavirus (2019-nCoV) Infection

Liana Zucco^{1, 2}, Nadav Levy^{1,2}, Desire Ketchandji³, Mike Aziz³, Satya Krishna Ramachandran¹ 1. Beth Israel Deaconess Medical Center Dept Anesthesia, Critical Care & Pain Medicine, Boston, USA 2. Healthcare Quality and Safety (MHQS), Harvard Medical School, Boston, USA 3. Oregon Health & Science University, Department of Anesthesiology & Perioperative Medicine, Portland, Oregon, USA

General

Your personal protection is the priority. Personal protective equipment (PPE) should be available for all providers to ensure droplet/contact isolation precautions can be achieved. Providers and organizations should review protocols for donning and doffing PPE. Careful attention is required to avoid self-contamination.

Patients with confirmed or suspected 2019-nCoV infected cases:

- Should NOT be brought to holding or PACU areas
- Should be managed in a designated OR, with signs posted on the doors to minimize staff exposure. Should be recovered in the OR or transferred to ICU into a negative pressure room. Ensure a high guality HMEF (Heat and Moisture Exchanging Filter) rated to remove at least 99.97% of airborne
- particles 0.3 microns or greater is placed between the ETT and reservoir bag during transfers to avoid contaminating the atmosphere.

Plan ahead:

o For time to allow all staff to apply PPE and barrier precautions o Consider intubation early to avoid the risk of a crash intubation when PPE cannot be applied safely.

During Airway Manipulation

- o Disposable mask, goggles, footwear, gown and gloves. Consider adopting the double glove technique.
- Standard ASA monitoring should be applied before induction of anesthesia. N95 mask at a minimum should be utilized. PAPR devices may offer superior protection when manipulating an airway of an infected patient.

Assign:

o Designate the most experienced anesthesia professionals available to perform intubation, if possible. Avoid trainee intubation for sick patients.

Avoid

o Awake fiberoptic intubation, unless specifically indicated. Atomized local anesthetic can aerosolize the virus.

Prepare to:

- Preoxygenate for 5 minutes with 100% FiO2
- o Perform a rapid sequence induction (RSI) to avoid manual ventilation of patient's lungs and potential aerosolization of virus from airways.
- Consider using a video-laryngoscope.

-200

 Depending on the clinical condition, the RSI may need to be modified. If manual ventilation is required, apply small tidal volumes

Use o Ensure there is a high quality HMEF (Heat and Moisture Exchanging Filter) rated to remove at least 99.97% of airborne particles 0.3 microns or greater placed in between the facemask and breathing circuit or between facemask and reservoir bag.

Dispose

 Re-sheath the laryngoscope immediately post intubation (double glove technique) Seal all used airway equipment in a double zip-locked plastic bag. It must then be removed for decontamination and disinfection.

Remember

o After removing protective equipment, avoid touching your hair or face before washing hands



Adapted from Kamming D, Gardam M, Chung F. I. Anaesthesia and SARS. Br J Anaesth 2003;90:715-18

- RSI when possible, 5 minutes of pre-oxygenation
- Video Laryngoscopy when possible
- Airborne Precautions, Contact, and Eye protection
- Double Glove
- Dispose of AIRWAY equipment properly





Medical Treatments

Respiratory Treatments for Hypoxia and ARDS

WORKING UP THE COVID19 PATIENT

Diagnostics

Common Lab Findings:



- Total lymphocyte count
- LDH, CRP, Ferritin, D-Dimer
- Procalcitonin
- AST/ALT

Chest Imaging: CXR vs CT vs US



Definitive Testing with RT-PCR





COVID 19 TREATMENT GUIDELINES FROM MASS GENERAL HOSPITAL/HARVARD

<u>fable 1:</u> Laboratories for diagnosis, prognosis / risk stratification, and/or safety of agents Suggested for <u>all hospitalized</u> patients with confirmed or suspected COVID-19					
Recommended daily labs:	<u>Viral serologies</u> : ²				
 CBC with diff (trend total lymphocyte count) Complete metabolic panel¹ CDV (i line) 	 HBV serologies (sAb, cAb, and sAg) HCV antibody, unless positive in past HUV 1/2 Ab/A a 				
CPK (creatine kinase) For risk stratification (may be repeated q2-3	HIV 1/2 Ab/Ag <u>If clinically indicated</u> :				
 <u>days if abnormal or with clinical deterioration</u>): D-dimer 	 Routine blood cultures (2 sets) For acute kidney injury (i.e. serum creatinine 				
• Ferritin / CRP / ESR	 For acute kiney injury (i.e. setum creatinine >0.3 above baseline), send urinalysis and spot urine protein:creatinine 				
 LDH Troponin³ 	<u>Procalcitonin</u>				
Baseline ECG ⁴ Radiology:	IL-6 See below for criteria Following up-to-date infection control				
 Portable CXR at admission High threshold for PA/lateral in ambulatory patients, consider only if low suspicion for 	 guidelines and appropriate PPE: SARS-CoV-2 test, if not already performed.⁵ If available, send influenza A/B and RSV test 				

patients, consider only if low suspicion for COVID-19 and result would change

CO-INFECTION IS RARE (LESS THAN 2%), **OBTAIN CORRECT LABS/DIAGNOSTICS**

Table 2: Risk Factors for Severe COVID-19 Disease					
Epidemiological – Category 1	Vital Signs – Category 2	Labs – Category 3			
Age > 55	Respiratory rate > 24	D-dimer > 1000 ng/mL			
	breaths/min				
Pre-existing pulmonary	Heart rate > 125 beats/min	CPK > twice upper limit of			
disease		normal			
Chronic kidney disease	SpO2 < 90% on ambient air	CRP > 100			
Diabetes with $A1c > 7.6\%$		LDH > 245 U/L			
History of hypertension		Elevated troponin			
History of cardiovascular		Admission absolute			
disease		lymphocyte count < 0.8			
Use of biologics		Ferritin > 300 ug/L			
History of transplant or other					
immunosuppression					
All patients with HIV					
(regardless of CD4 count)					

RISK CATEGORIZE PATIENTS



AVOID STEROIDS AND NSAIDS

 NSAIDS link to COVID19 unclear, however concern that it may worsen outcomes

- may increase ACE receptor expression?
- Steroids should be avoided unless absolutely necessary

ADVANCED TECHNIQUES FOR HYPOXIA AND ARDS

AVOID AERSOLIZATION

If possible,

No Nebulized Treatments

No Bag Mask Ventilation

No Heated High Flow Nasal Cannula

No Non-Invasive Ventilation (e.g. BiPAP)

Use a HEPA Filter

- Data from China shows that early intubation is preferred based on Disease progression Characteristics
- Anyone with a 6L requirement or greater should be transferred to an ICU and intubated (Current Cedars Sinai Protocol)
- BIPAP and High Flow Nasal cannula Not advised
- Days on Mechanical ventilation 6 to 10

EARLY INTUBATION!

An acute condition characterized by bilateral pulmonary infiltrates and severe hypoxemia in the absence of evidence for cardiogenic pulmonary edema.

- 1. Acute onset
- 2. Bilateral disease
- 3. Hypoxemia
- 4. Non-cardiogenic

Basic Criteria



ACUTE RESPIRATORY DISTRESS SYNDROME





P:F Ratio

The P:F Ratio is calculated by: PaO2/FiO2 BERLIN CRITERIA : Mild ARDS = P/F <300 but >200 Moderate ARDS = P/F <200 but >100 Severe ARDS = P/F <100

Figure 6-2. University of Michigan ARDS algorithm: sample stepwise approach to mechanical ventilation in acute respiratory distress syndrome (ARDS)



1: Basic LPVS ARDS Network ventilation strategy: a. Using VCV or PCV and targeting VT 4-6 b. Maintain Pplat <30 cm H₂O c. PEEP/FiO₂ per table (see back page) 2: Pt-Vent Dyssynchrony, Step 1 Initial strategy should be to: a. Assess potential to treat with pharmacologic agents (eg. sedation, NMB agents), and b. Consider minor ventilator adjustments (eg. flow rate & pattern, inspiratory pause) If above does not work, consider increasing VT 1 mL/kg (max 8 mL/kg), provided Pplat 3: Pt-Vent Dyssynchrony, Step 2 Consider a variable flow pressure breath mode b. Pressure Control Ventilation 4: Criteria for Failing LPVS PaO₂ <55 torr on FiO₂ =1.0 and Pplat >30 cm H₂O on VT =4 mL/kg PBW Use unit specific rotation frequency, but evidence suggests majority Recommend a 48 hr trial, stop if no improvement, as evidenced by: Pressure Control Inversed Ratio (PCIRV) Ventilation Airway Pressure Release Ventilation (APRV) Esophageal Pressure (Pes) Guided Therapy High Frequency Oscillatory Ventilation (HFOV) If positive response, titrate and consider iloprost, per Respiratory Extracorporal Membrane Oxygenation (ECMO) Absolute contraindications: irreversible pulmonary process and Evaluate, but lower survival if on vent 7-10 days pre-ECMO

Evidenced Based Strategies for Reducing Mortality

- Lung Protective Ventilation (ARDSNET)
- Paralysis (ACURASYS/ROSE-PETAL)
- PRONING (PROSEVA)
- Conservative Fluid Resuscitation (FACCT)
- ECMO (CESAR and EOLIA)

> Access to Ventilators contributes reduction in Mortality

University of Michigan: <u>https://aneskey.com/on-</u> acute-respiratory-distress-syndrome/

Lung Protective Ventilation

ARDS NETWORK 2000

Reduce volu/barotrauma

6 mL/kg Predicted Body Weight

Recommend starting at 8mL/kg and titrating down as able, adjust RR to maintain MV

If Pplat > 30 cmH20 at 6mL/kg, decreased VT as able to achieve Pplat < 30 cmH2) (minimum of 4mL/kg)

Significantly improves mortality and decreases ventilator days



NIH NHLBI ARDS Clinical Network

Disadvantages: Uncomfortable Decreases ventilation

PEEP Ladder

ARDS NETWORK 2000

FiO2	0.3	0.4	0.5	0.6	0.7	0.8	0.9	1.00
PEEP	5	5-8	8-10	10	10-14	14	14-18	18-24

Oxygenation Goal: PaO2 55-80mmHg or SpO2 88-95%

Arter al	1			
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2.1	%	[- 1.5
12.7	%	[- 5.0
0.3	%	[0.0	- 1.5
				and the second
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131	mmol/L			
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96	mmol/L]	94	- 107
				Statist.
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1.2	mmol/L	1	0.5	- 2.0
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				and the
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Permissive Hypercapnia

Smaller VT often means decreased ventilation

PaCO2=65mmHg and/or pH=7.20 generally very well tolerated

Sodium bicarbonate infusion generally not recommended

Hypercapnia MAY confer physiologic benefit

and Acute Respiratory Distress Syndrome, UT SOUTHWESTERN



Agarwal, S et al. Prone position ventilation in ARDS. Indian Journal of Anesthesia, Vol 59, No.4, April, 2015, 246-248.

Prone Positioning (PROSEVA 2013)

Prone positioning confers physiologic benefit by several mechanisms:

Improves V/Q Matching

Decreases lung compression

Improves bronchial drainage

Potentially increases FRC

51% relative mortality risk reduction with early proning



Guerin C, et al. PROSEVA. NEJM 2013

ACURASYS – THE PREVIOUS STUDY



- Landmark Study, Multi-center, Randomized Controlled Trial.
- Significant mortality improvement in the Cisatricurium group in patients with ARDS
- No standardization for proning, steroids, nitric oxide use

(ACURYSYS) Papazian L, Forel J-M, Gacouin A, et al. Neuromuscular blockers in early acute respiratory distress syndrome. N Engl J Med 2010;363:1107-1116

- > AKA "Rose-Petal" Trial, Randomized, Multi-Center
- Inclusion Criteria
 - ► P/F <150mmHg
 - ► PEEP>8
 - Bilateral Opacities
 - Not pulmonary edema/fluid Overload
- Exclusion Criteria
 - ► ECMO
 - > NMB prior to enrollment
 - > P:F> 200 at time of randomization.

(ROSE) National Heart, Lung, and Blood Institute PETAL Clinical Trials Network Moss M, Huang DT, et al. Early Neuromuscular Blockade in the Acute Respiratory Distress Syndrome. N Engl J Med 2019; 380:1997.

ORIGINAL ARTICLE

Early Neuromuscular Blockade in the Acute Respiratory Distress Syndrome

The National Heart, Lung, and Blood Institute PETAL Clinical Trials Network*

Article Figures/Media

Metrics

May 23, 2019 N Engl J Med 2019; 380:1997-2008 DOI: 10.1056/NEJMoa1901686

43 References 5 Citing Articles 3 Comments

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- Randomly assigned patients with moderate to severe ARDS (PaO2 less than 150, Peep > 8cm) to Cisatracurium or "usual-care approach" without routine NMB blockade.
- > End point was in-hospital death at 90 days
- Larger trial, very strict protocol, Use of higher PEEP (5 in Acurasys)
- Lighter Sedation Targets in non-intervention group (Possible confounder)
- Exclusion criteria possibly Achilles heel (655 received neuromuscular blockade)
- Large difference between proning between this (fewer proned patients) and ACURASYS.

KOSE) National Heart, Lung, and Blood Institute PETAL Clinical Trials Network, Moss M, Huang DT, et al. Early Neuromuscular Blockade in the Acute Respiratory Distress Syndrome. N Engl J Med 2019; 380:1997.

- For severe ventilator dis-synchrony I will continue to use paralysis
- > NMB still acceptable, shows no harm
- Lack of benefit, may be because underpowered for the study population
 - Vent Dis-synchrony may not have been adequately addressed and significant proportion of already paralyzed patients excluded

MY TAKE ON PARALYSIS (ACURASYS VS ROSE-PETAL)



Source: Tobin MJ: Principles and Practice of Mechanical Ventilation, 3rd Edition: www.accessanesthesiology.com

Pulmonary Vasodilators

Examples: Epoprostenol, inhaled nitric oxide

Improves P:F ratio, at least transiently

No significant benefit with regard to mortality or duration of mechanical ventilation

Potential adverse effects include decreased platelet aggregation, renal dysfunction and methemoglobinemia

ORIGINAL ARTICLE

Comparison of Two Fluid-Management Strategies in Acute Lung Injury

The National Heart, Lung, and Blood Institute Acute Respiratory Distress Syndrome (ARDS) Clinical Trials Network*

ABSTRACT

BACKGROUND

Optimal fluid management in patients with acute lung injury is unknown. Diuresis or fluid restriction may improve lung function but could jeopardize extrapulmonaryorgan perfusion.

METHODS

In a randomized study, we compared a conservative and a liberal strategy of fluid management using explicit protocols applied for seven days in 1000 patients with acute lung injury. The primary end point was death at 60 days. Secondary end points included the number of ventilator-free days and organ-failure-free days and measures of lung physiology.

RESULTS

The rate of death at 60 days was 25.5 percent in the conservative-strategy group and 28.4 percent in the liberal-strategy group (P=0.30; 95 percent confidence interval for the difference, -2.6 to 8.4 percent). The mean (±SE) cumulative fluid balance during the first seven days was -136±491 ml in the conservative-strategy group and 6992±502 ml in the liberal-strategy group (P<0.001). As compared with the liberal strategy, the conservative strategy improved the oxygenation index (Imean airway pressure × the ratio of the fraction of inspired oxygen to the partial pressure of arterial oxygen] × 1000 and the lung injury score and increased the number of ventilator-free days (14.6±0.5 vs. 12.1±0.5, P<0.001) and days not spent in the intensive care unit (13.4±0.4 vs. 11.2±0.4, P<0.001) during the first 28 days but did not increase the incidence or prevalence of shock during the study or the use of dialysis during the first 60 days (10 percent vs. 14 percent, P=0.06).

CONCLUSIONS

Although there was no significant difference in the primary outcome of 60-day mortality, the conservative strategy of fluid management improved lung function and shortened the duration of mechanical ventilation and intensive care without increasing nonpulmonary-organ failures. These results support the use of a conservative strategy of fluid management in patients with acute lung injury. (ClinicalTrials. gov number, NCT00281268.)

Conservative Fluid Administration

FACTT 2006

Compared conservative vs. liberal fluid management strategies in patients with "ALI"

No difference in mortality at 60 days

Decreased ICU time and increased ventilator-free days in conservative group

- CESAR Trial (Lancet 2009): Decreased mortality and disability at 6 months for patients with severe ARDS transferred to ECMO center
- EOLIA Trial (NEJM 2018): No difference between early use of VV ECMO and standard MV with VV ECMO as rescue
- "WHO interim guidelines recommend offering extracorporeal membrane oxygenation (ECMO) to eligible patients with acute respiratory distress syndrome (ARDS) related to coronavirus disease 2019 (COVID-19). "

ECMO



A Seattle Intensivist's One-pager on COVID-19

Nomenclature

Infection: Coronavirus Disease 2019 a.k.a. COVID-19 Virus: SARS-CoV-2, 2019 Novel Coronavirus NOT "Wuhan Virus"

Biology

- 30 kbp, +ssRNA, enveloped coronavirus
- Likely zoonotic infection; source/reservoir unclear (Bats? / Pangolins? → people)
- Now spread primarily person to person;
 - Can be spread by asymptomatic carriers!
- Viral particles enter into lungs via droplets
- Viral S spike binds to ACE2 on type two pneumocytes
- Effect of ACE/ARB is unclear; ACE vs ARBs may even have opposite effects
- Other routes of infection (contact, enteric) possible but unclear if these are significant means of spread

Epidemiology

- Attack rate = 30-40%
- $R_0 = 2-4$ (similar to influenza)
- CFR = 3.4% (worldwide numbers)
- Incubation time = 4-14 days typically (up to 24 days) Timeline:
- China notifies WHO 2019-12-31
- First US case in Seattle 2020-1-15
- WHO Declared pandemic 2020-3-11
- National emergency 2020-3-12

Disease clusters: SNFs, Conferences, other Strategies: contact tracing, screening, social distancing



Diagnosis/Presentation

- Symptoms
- 65-80% cough ٠
- 45% febrile on presentation (85% febrile during illness)
- 20-40% dyspnea
- 15% URI symptoms
- 10% GI symptoms

Labs

- CBC: Leukopenia & lymphopenia (80%+) ٠
- BMP: **↑**BUN/Cr ٠
- LFTs: ↑AST/ALT/Tbili
- ↑ D-dimer, ↑ CRP, ↑ LDH
- ↑ IL-6, ↑ Ferritin ٠
- ↓ Procalcitonin
- *PCT may be high w/ bacterial superinfxn*

Imagine

- . CXR: hazy bilateral, peripheral opacities
- CT: ground glass opacities (GGO), crazy paving, consolidation, *rarely may be unilateral*



- POCUS: numerous B-lines, pleural line thickening, consolidations w/ air bronchograms Isolation
- Phone call is the best isolation (e.g. move to telemed)
- Place patient in mask, single room, limit/restrict visitors -

Precautions

- STANDARD + CONTACT (double glove) +
- Either AIRBORNE (for aerosolizing procedures: intubation, extubation, NIPPV, suctioning, etc) or DROPLET (for everything else)
- -N95 masks must be fit tested; wear eye protection
- PPE should be donned/doffed with trained observer
- Hand hygiene: 20+ seconds w/ soap/water or alcohol containing hand gel

Treatment

TBil

DBili

BUN CI \uparrow

Cr

нсоз 🛧

Glu

Alk

Phos \uparrow

ALT

Na

- Isolate & send PCR test early (may take days to result)
- GOC discussion / triage
- Notify DOH, CDC, etc
- Fluid sparing resuscitation
- ± empiric antibiotics
- Intubate early under controlled conditions if possible
- Avoid HFNC or NIPPV (aerosolizes virus) unless individualized reasons exist (e.g. COPD, DNI status, etc); consider helmet mask interface (if available) if using NIPPV
- Mechanical ventilation for ARDS
 - LPV per ARDSnet protocol
 - 7 P's for good care of ARDS patients: e.g. PEEP/Paralytics/Proning/inhaled Prostacyclins, etc
 - ? High PEEP ladder may be better
 - ? ECMO in select cases (unclear who)
- Consider using POCUS to monitor/evaluate lungs
- Investigational therapies:
 - Remdesivir -- | block RNA dependent polymerase
 - Chloroquine -- | blocks viral entry in endosome
 - Tocilizumab -- | block IL-6
 - Corticosteroids -- | reduce inflammation
- None of these investigational therapies are proven, but literature is evolving quickly.

Prognosis

Age and comorbidities (DM, COPD, CVD) are significant predictors of poor clinical outcome; admission SOFA score also predicts mortality.

14%

- Lab findings also predict mortality
 - ↑ d-dimer,

 - ↑ cardiac
 - myoglobin
- Expect prolonged MV
- Watch for complications: Secondary infection (VAP), Stress CM, etc





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- SCCM Resource availability https://sccm.org/Blog/March-2020/United-States-Resource-Availability-for-COVID-19?_zs=jxpjd1&_zl=w9pb6
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- ▶ Feel free to reach out.
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