Managing Patients in the Transition Between Floor and ICU

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Disclosures

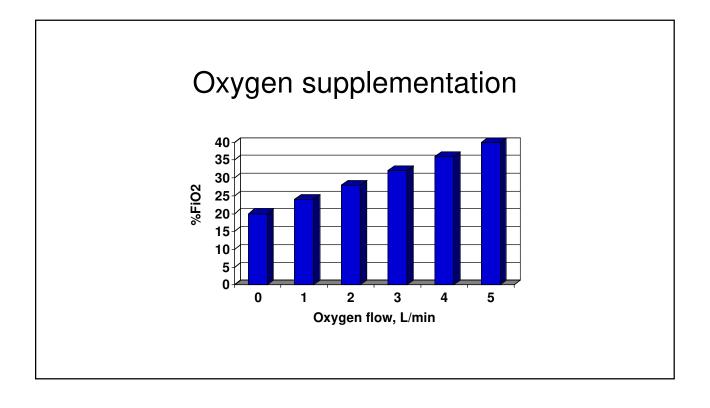
• None

- 37 M mild asthma, type 2 diabetes, morbid obesity, recent new smoker
- Presents with 5 days of productive cough, fever, progressive dyspnea
- Exam: T 38.7 °C (101.7 °F), RR 20, Sat 90% on room air, bibasilar crackles, mild expiratory wheezing
- WBC 16, Na 131
- ABG on room air: pH 7.31, PaCO₂ 52, PaO₂ 63
- CXR: bilateral infiltrates
- CT chest: ground-glass opacities, worse in bases and periphery

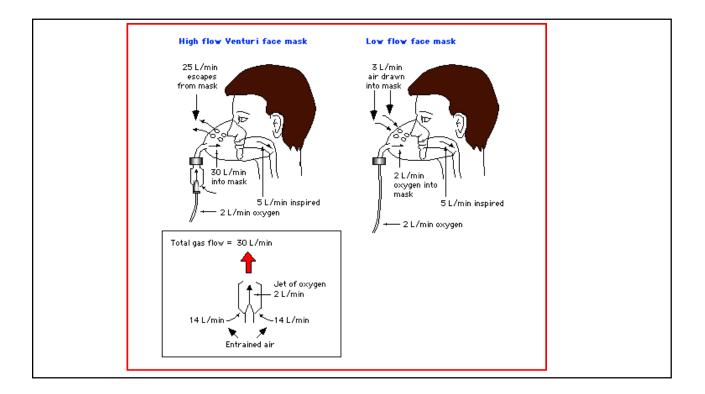


Case 1: Imaging

- Started on ceftriaxone/azithro for presumed communityacquired pneumonia, bronchodilators
- On hospital day 2, respiratory status is worse
- RR 30, on 50% face mask
- ABG: pH 7.46, PaCO₂ 33, PaO2 66
- What options can be used to increase respiratory support on the floor?



Method	FiO2	Flowrate	
	(Approximate)	(L/min)	
Non rebreather Mask	60-80%	10-15	
Venti Mask	24%	3	
	26%	3	
	28%	6	
	31%	6	
	35%	9	
	40%	12	
	50%	15	
Simple Face Mask	35-55%	5-10 lpm	
Nasal Cannula	24%	1	
	28%	2	
	32%	3	
	36%	4	
	40%	5	
	44%	6	

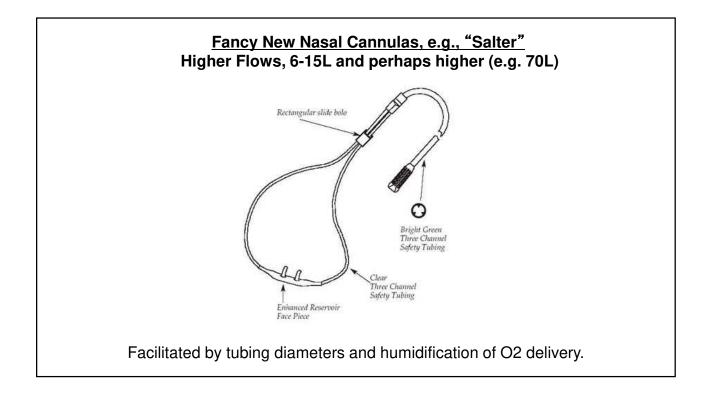


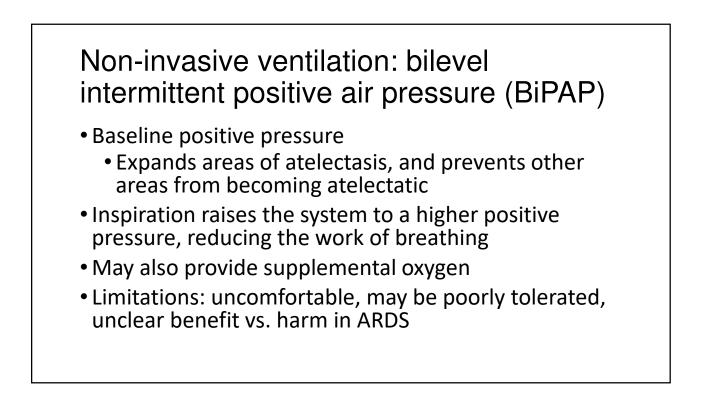
Case 1 continued

- Increasing O2 requirements to 10L
- Continued desaturations and increased work of breathing
- Next steps?
- High flow NC vs NIPPV?

High-flow nasal cannula (HFNC)

- 30 to 60 LPM oxygen heated to body temperature at 100% relative humidity
- Dramatically reduces dilution of inspired oxygen by room air
- · Washes out dead space in the airways
- Can generate low-level continuous positive airway pressure (CPAP) at higher flows
- · Moistens secretions, improves mucociliary transport
- Greater comfort and adherence
- Limitations: doesn't provide as much positive pressure or reduce the work of breathing as much as non-invasive ventilation; does it help prevent intubation, or "delay the inevitable"?





Noninvasive positive pressure ventilation (NIPPV)

- Supportive data for:
 - Acute COPD exacerbation--greatest benefit
 - Ventilator weaning adjunct in COPD
 - Acute cardiogenic pulmonary edema
 - Hypoxemic respiratory failure, immunocompromised host in early but not later studies
- Why?
 - Improved alveolar ventilation
 - Reduced work of breathing
 - Relieving fatigued respiratory muscles
- Outcomes
 - Decreased infections
 - Fewer intubations

NIPPV, cont'd

- Facemask: better effect, less comfort
- Need "protocol-driven" initiation, with careful monitoring, esp in 1st 1-2 hours
- Contraindications:
 - Mental status, hemodynamics, facial deformity, upper airway obstruction, SECRETIONS, aspiration risk
 - Initial concern re: aerosolization in COVID; unclear benefit vs. harm in lung injury

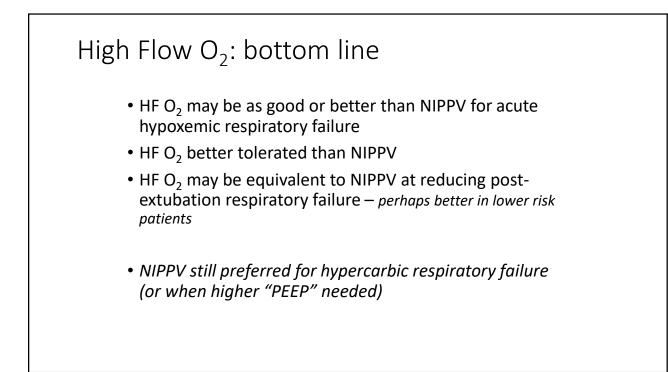


HFNC vs BiPAP in clinical trials

In a meta-analysis of 29 RCTs, HFNC was associated with lower need for intubation, lower mortality, and greater comfort than non-invasive ventilation

ACP guidelines lean toward HFNC as the preference over NIV for acute AHRF and over conventional oxygen for postextubation AHRF with "low-certainty evidence"; (my own editorial: NIV has better rationale than HFNC for more significant *hypercapnic* respiratory failure)

Ann Intern Med 2021;174:952



Case 1 continued

- Started on high flow NC
- Ongoing desaturations and escalating O2 requirement
- Anesthesia called for intubation

What is the optimal timing for intubation in the patient with deteriorating respiratory status?

As an example: 2019 AHA update: Airways

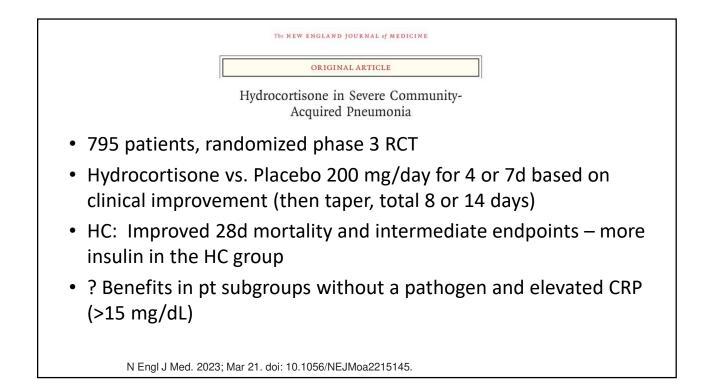
- Data isn't clear when/how to intubate during a code situation due to varying patient and provider circumstances.
- Use best judgement based on situation and provider expertise.
- It is suggested that proficiency be encouraged to acquire and maintain airway insertion skills for relevant personnel.

Case 1 evolution

A bronchoscopy was performed with cell count/differential:
– 685 RBC, 295 WBC (32% Neutrophils, 13% Lymphs, 8% Monos, 39% eosinophils)

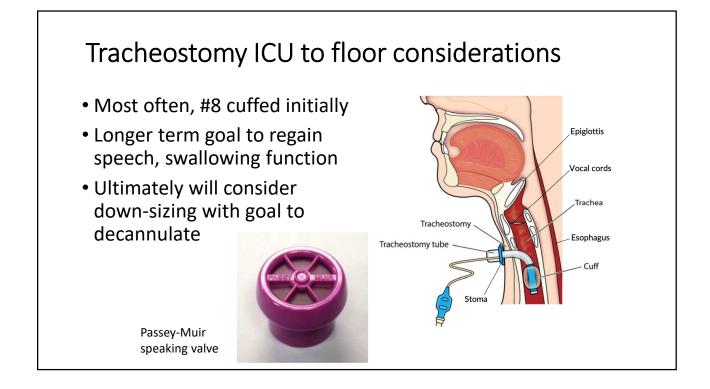
Acute Eosinophilic pneumonia

- Known causes: e.g., meds, parasites, new onset smoking
- DDx: "PIE", e.g., Churg Strauss, CEP, ABPA, etc
- Idiopathic:
 - Similar presentation as CAP
 - Classically no circulating eosinophils detected, but eos seen in BAL
 - Can respond dramatically to steroids
 - One approach is to treat for 1 month after symptoms disappear and CXR normal (i.e., on the order of 4-6 weeks); inhaled steroids may be beneficial in relapse



Case 1 conclusion

- Intubated for 2 weeks with slow improvement
- Required trach
- Discharged to a ventilator weaning rehab facility

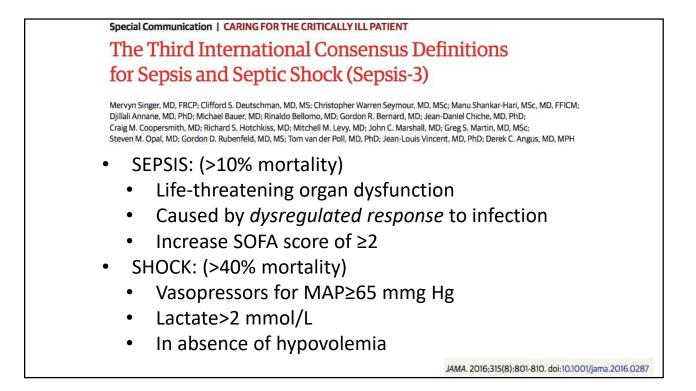


- 87 F PMHx type 2 diabetes on metformin
- Brought in with confusion, hypoglycemia
- Given D50W in the field by EMS for glucose 40 mg/dl (2.2 mmol/L)
- Exam: T 92°F (33.3°C), P 90, BP 65/40, RR 20, Sats 97% RA

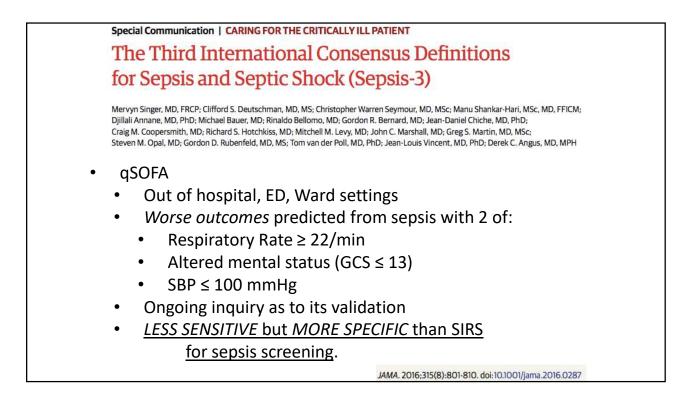
- Laboratory workup:
- WBC 18, Na 124, K 5.2, Bicarb 11, BUN 60 mg/dl, Cr 4.0 mg/dl (354 μmol/L)
- Urinalysis >100 WBC, 4+ bacteria
- Is this sepsis? Next steps?

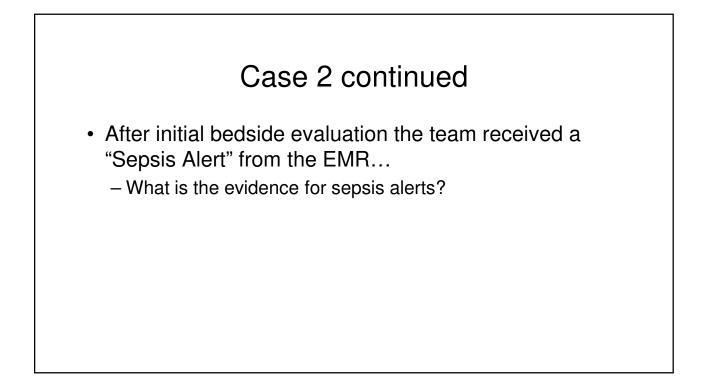
What is Sepsis (2001-15)?

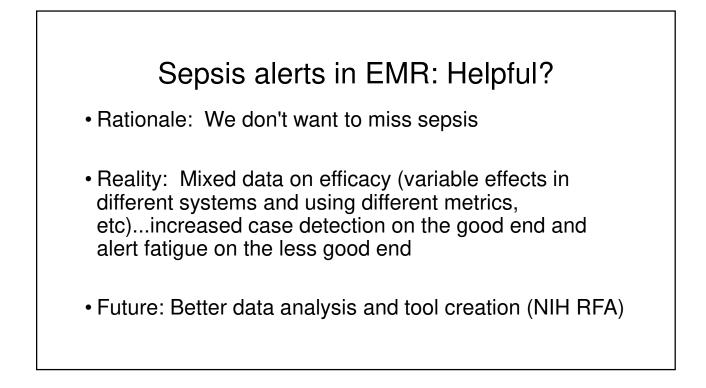
- <u>Systemic Inflammatory</u> <u>Response Syndrome</u> (SIRS):
 - Temp >38°C or <36°C
 - Heart Rate > 90 bpm
 - Resp Rate > 20/min
 - WBC >10000, <4000, or Bandemia>10%
- <u>Sepsis</u>: SIRS + Infection
- <u>Severe Sepsis</u>: Sepsis+ Organ Dysfunction
- <u>Septic Shock</u>: Sepsis+Refractory Hypotension

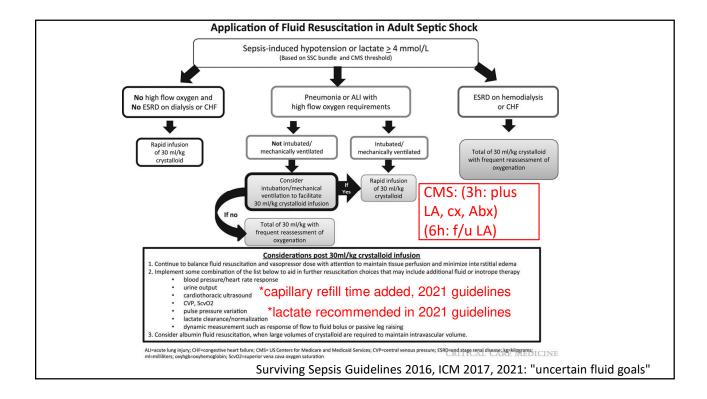


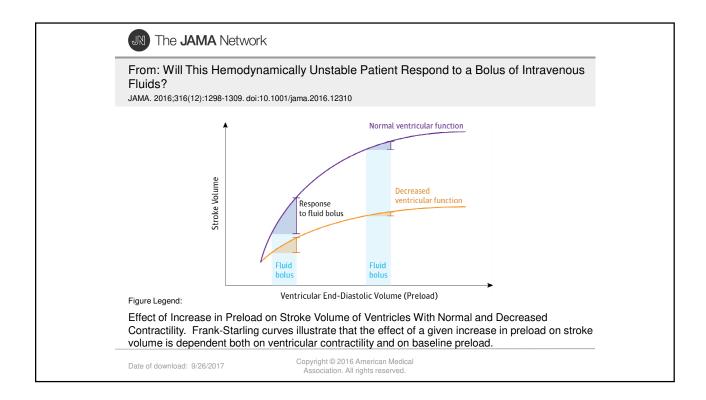
Points	0	1	2	3	4
PaO2/FiO2	≥400	<400	<300	<200 + mechanical ventilation	<100 + mechanical ventilation
Platelets	≥150	<150	<100	<50	<20
Bilirubin	<1.2	1.2-1.9	2.0-5.9	6.0-11.9	>12.0
Blood Pressure	MAP ≥70	MAP <70	Dopamine <5 or Dobutamine	Dopamine 5.1-15 or Epinephrine <0.1 or Norepinephrine <0.1	Dopamine >15 or Epinephrine >0.1 or Norepinephrine >0.1
Glasgow Coma Scale	15	13-14	10-12	6-9	<6
Creatinine	<1.2	1.2-1.9	2.0-3.4	3.5-4.9 or <500cc urine/d	>5.0 or <200cc urine/d

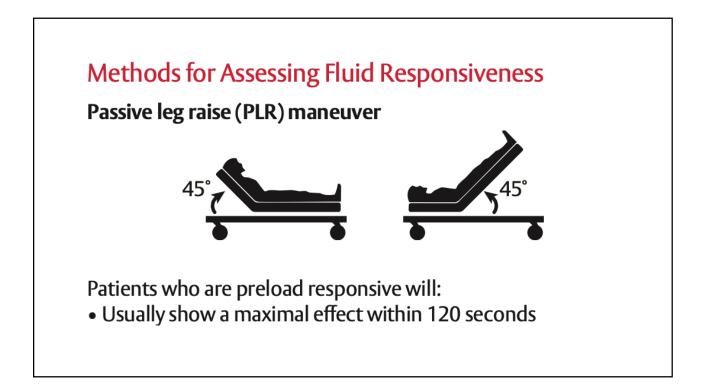


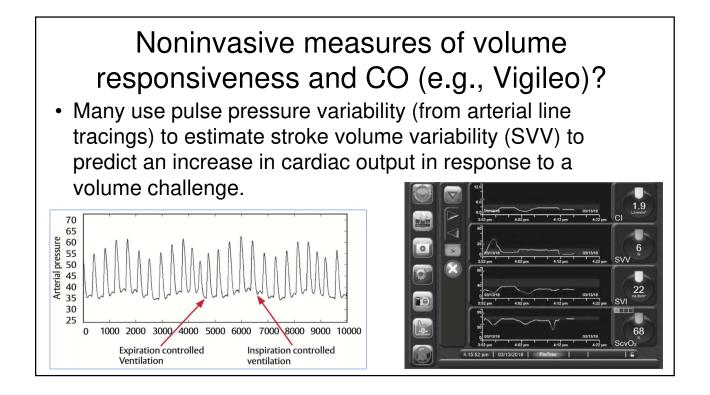


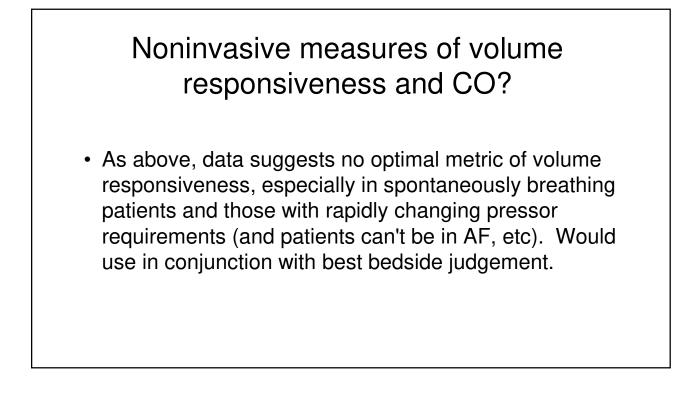


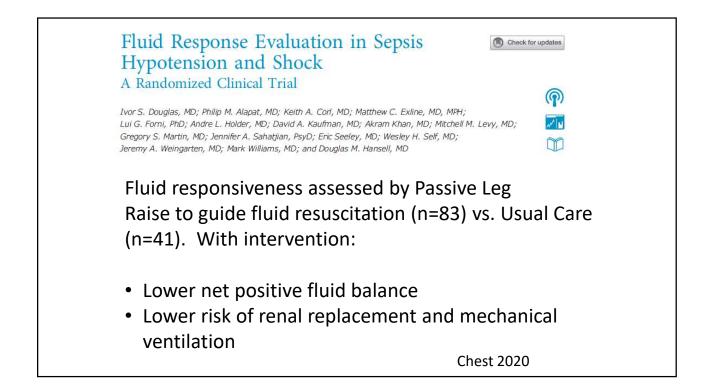


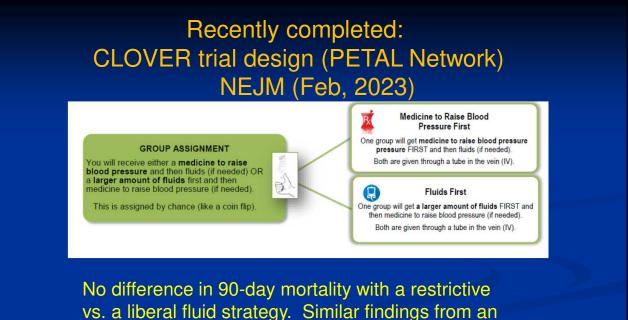




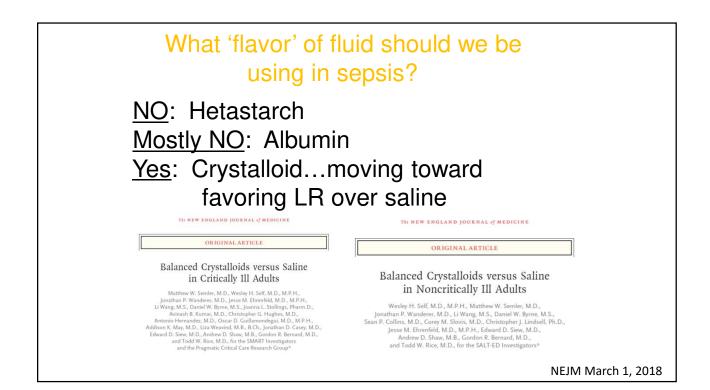


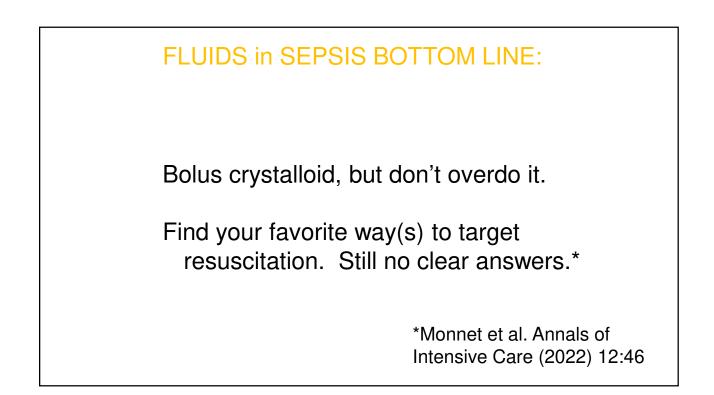






Denmark study (NEJM June, 2022).





Other issues related to sepsis, including other 2021 guidelines points

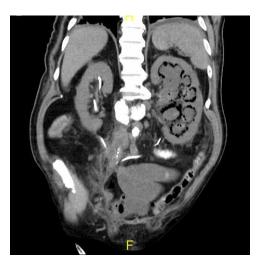
- Optimal antibiotics dosing
- Acknowledges reasonable to dose pressors peripherally while awaiting central access
- Long term follow-up needs in critical illness survivors
- Accumulating data is not supporting a role for Vitamin C in sepsis (NEJM June 2022)

Case 2 continued

- Ongoing hypotension despite adequate fluid resuscitation
- Started on norepinephrine through peripheral IV
- Transferred to ICU for further management

Case 2 continued

- Continued instability led to urgent imaging
- Emphysematous pyelonephritis
- Necrotizing soft tissue infection
- Often underlying DM
- · May be associated with obstruction
- Treatment: antibiotics and/or drainage; surgery in severe cases where feasible



Case 2 continued

- Cultures grew E. coli; slowly responded to antibiotics no obstruction on imaging and poor surgical candidate, regardless.
- Course complicated by AKI requiring HD, delirium
- Hemodynamically stable but still requiring a lot of care:
 - Delirious
 - Extremely weak
 - End of life and goals of care discussions with family have been challenging
- Transfer to floor!

Take Home Messages

- Wide spectrum of illness from floor to ICU and back
- Multiple options exist for O2 delivery and respiratory support
- Early recognition and treatment of sepsis is key
- Judicious fluid resuscitation (avoiding volume overload) is important
- Close communication between floor and ICU is critical for optimal management