

Antibiotic Recommendations for Sepsis and Septic Shock

- This guidance is for patients with sepsis and septic shock (Sepsis 3 criteria) **only** where early initiation of active antibiotics has been shown to improve outcomes
- Those with less severe infections should have syndromic antibiotics started per NM guidelines available via the stewardship website and/or One Chart order sets
- Appropriate cultures should be obtained which include two sets of blood cultures obtained before antibiotics are started and cultures of other suspected sites of infection (sputum, urine, etc.) obtained as soon as possible
- Use of two antibiotics targeting gram negative pathogens (i.e. combination therapy) is **not** routinely recommended, but may be considered in patients with septic shock
 - If two agents are started, they should rapidly be narrowed when culture becomes available
- Antibiotics started for sepsis should be narrowed to target pathogens isolated as soon as culture results become available or clinical improvement is achieved in the absence of culture data
 - Antibiotics started for sepsis should be reassessed daily for potential discontinuing if infection is ruled out or narrowing if more data becomes available
 - Patients with consistently low procalcitonin values (<0.5) can usually have antibiotics safely stopped

Suspected Source of Infection	Antibiotic Recommendations
Unknown Source (includes catheter related blood stream infection) +	<p>Vancomycin IV* PLUS Cefepime 1 gm IV q6hr +/- Tobramycin 7 mg/kg IV EIAD⁺</p> <p><u>Severe beta-lactam allergy (anaphylaxis, hives):</u> Vancomycin IV PLUS Aztreonam 2g q8h +/- Tobramycin 7 mg/kg IV EIAD⁺</p> <p>+ Consider addition of micafungin 100mg daily in those at high risk for candidemia. Risk factors for candidemia at NM include: 1) Broad-spectrum antibiotic use, 2) Central venous catheter, 3) Receipt of TPN, 4) Recent abdominal surgery, and 5) Steroid use. Presence of 2 or fewer of the risk factors suggests a 99.4% chance of not developing candidemia, while patients with >2 risk factors have a 4.7% risk of developing candidemia.</p>
Intra-abdominal Source	<p>Piperacillin/tazobactam 4.5g IV q8h, over 4 hours OR Cefepime 1g q6h hours PLUS Metronidazole 500 mg IV q8h +/- Gentamicin 7 mg/kg IV EIAD</p>

<p>Severe Community Acquired Pneumonia – No Risk Factors for resistance (see below for risk factors)</p> <p><u>Risk Factors for MRSA</u> = Documented MRSA sputum colonization, post-influenza pneumonia, severe necrotizing pneumonia</p> <p><u>Risk Factors for resistant gram-negative rods</u> = history of sputum colonization with Pseudomonas or organisms resistant to typical CAP therapy</p> <p><u>Risk Factors for both MRSA and Resistant Gram-negative rods</u> = Recent hospital stay with use of IV antibiotics (>5 days)</p>	<p>Ceftriaxone 2 gm IV q24h⁺ PLUS Azithromycin 500 mg IV q24h</p> <p><u>Severe beta-lactam allergy (anaphylaxis, hives):</u> Levofloxacin 500 mg IV q24h⁺</p> <p><u>MRSA Risk:</u> Consider addition of Vancomycin IV* or Linezolid to above</p> <p><u>Resistant Gram-negative Rod Risk:</u> Consider Piperacillin/tazobactam PLUS Azithromycin OR Cefepime PLUS Azithromycin</p> <p><u>Risk Factors for MRSA and Resistant Gram-negative Rods:</u> Consider Vancomycin IV* PLUS Cefepime PLUS Azithromycin</p>
<p>Nosocomial Pneumonia Hospital-acquired pneumonia (HAP) and ventilator-associated pneumonia (VAP)</p> <p>Treat using CAP guidelines if hospitalized ≤5 days</p>	<p>Vancomycin IV* PLUS Cefepime 1 gm IV q6hr +/- Tobramycin 7 mg/kg IV EIAD (if concern for Pseudomonas)</p> <p><u>Severe beta-lactam allergy (anaphylaxis, hives):</u> Vancomycin IV* PLUS Aztreonam 2g q8h +/- Tobramycin 7 mg/kg IV EIAD (if concern for Pseudomonas)</p>

* Vancomycin dosed per pharmacy consult. Typically with loaded with 20-25 mg/kg dose initially (max 2g initial dose)

EIAD: Extended Interval Aminoglycoside Dosing

Updated: August 2021