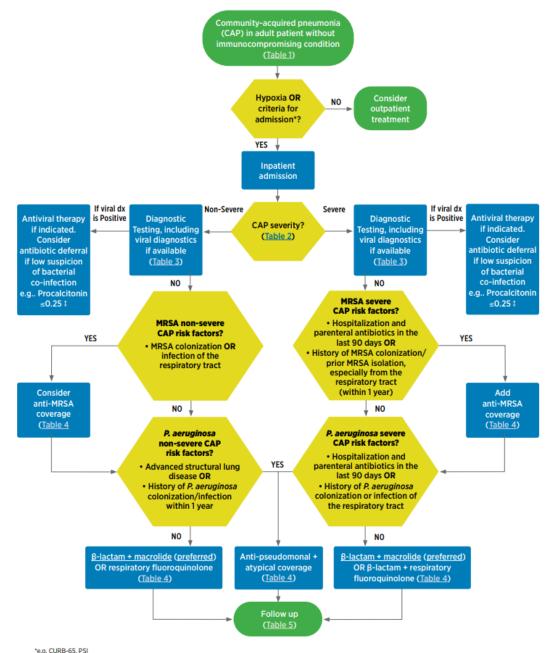


Stony Brook University Hospital Treatment Guidelines:

Management of Community Acquired Bacterial Pneumonia (CABP) in Adult Patients



*e.g. CURB-65, PSI ‡ This is a clinical practice enhancement to the ATS/IDSA CAP clinical practice guideline

 $\frac{https://www.idsociety.org/globalassets/idsa/practice-guidelines/community-acquired-pneumonia-in-adults/cap-clinical-pathway-final-online.pdf$

1. Background

1. SBUH has incorporated national guidelines in its creation of hospital-wide treatment guidelines for the management of CABP in adult patients to guide appropriate antibiotic use.

2. Definitions¹

- 1. <u>Pneumonia:</u> Infection of the pulmonary parenchyma characterized by the filling of the alveoli with fluid
 - i. Can result if there is a defect in the host's defense or if the host's defense is overcome by high inoculum or virulence
- 2. <u>Community-acquired bacterial pneumonia:</u> Pneumonia acquired either from the community setting or that occurs within 48 hours of a hospital admission, from a confirmed or presumed bacterial etiology

3. Symptoms²

1. Fever, chills, cough, dyspnea, pleuritic chest pain, headache, myalgia

4. Diagnosis²

- 1. Identification of infiltrate or effusion on chest radiograph or other imaging technique with clinical symptoms of pneumonia
- 2. Pre-treatment gram stains, respiratory secretion cultures, and blood cultures are recommended in the hospital setting only in certain situations
 - i. Severe CABP, especially if intubated
 - ii. Risk factors for methicillin-resistant S. aureus (MRSA) or P. aeruginosa

5. Risk Factors for MRSA or P. aeruginosa

- 1. Prior respiratory isolation of MRSA or *P. aeruginosa*
- 2. Recent hospitalization and parenteral antibiotics in the last 90 days

6. Severity Classification of CABP²

1. If patients do **not** meet these criteria, they are considered to have non-severe CABP

Table 1. Criteria for Defining Severe CABP

Either 1 Major Criterion OR ≥ 3 Minor Criteria

Major Criteria

- 1. Septic shock with need for vasopressors
- 2. Respiratory failure requiring mechanical ventilation

Minor Criteria

- 1. Respiratory rate ≥ 30 breaths/min
- 2. PaO2/FiO2 ≤ 250
- 3. Multilobar infiltrates
- 4. Confusion or disorientation
- 5. Uremia (BUN ≥ 20 mg/dL)
- 6. Leukopenia (WBC < $4,000 \text{ cells/}\mu\text{L}$)
- 7. Thrombocytopenia (platelets < 100,000/μL)
- 8. Hypothermia (temperature < 36°C)
- 9. Hypotension requiring aggressive fluid resuscitation

7. Criteria for IV to PO conversion

Table 2. Criteria for IV to PO conversion

Criteria for IV to PO Conversion

- 1. Heart rate < 100 beats/min
- 2. Respiratory rate < 20 breaths/min
- 3. Blood pressure stable
- 4. Temperature < 38°C for 24 hours
- 5. WBC < 15,000 cells/μL
- 6. Hemodynamically stable
- 7. Patient able to tolerate PO medications

8. Antibiotic Selection²

1. SBUH Antibiogram Data

- i. Please consult SBUH Antibiogram for institutional susceptibility patterns when selecting therapy (https://renaissance.stonybrookmedicine.edu/medicine/asp)
- 2. Table 3 lists treatment options for empiric therapy in adult patients with CABP.
 - Ceftriaxone with either azithromycin or doxycycline is the preferred treatment for nonsevere CABP
 - 1. Doxycycline may be used instead of azithromycin in persons with a known history of cardiac arrhythmia or prolonged QTc intervals on ECG
 - ii. Fluoroquinolone monotherapy may be considered in persons with suspected or confirmed *Legionella* infection
- 3. If patient has risk factors for (refer to Section 5) <u>OR</u> a known history of infection with MRSA or *P. aeruginosa*, please refer to **Table 4** for targeted treatment options
 - i. If MRSA surveillance results with no growth, MRSA coverage can be de-escalated
 - i. If microbiology results without isolation of these organisms, broad coverage can be deescalated as treatment regimen should be targeted based on microbiological results
- 4. Addition of anaerobic coverage (i.e. metronidazole) is **not recommended** for suspected aspiration pneumonia
- 5. Use of broad-spectrum antibiotic coverage such as meropenem, piperacillin-tazobactam, and cefepime is usually **not** needed for CABP treatment if the patient has recent healthcare system exposure (i.e. nursing home, hemodialysis)
 - Use of a validated clinical decision support tool such as the Drug Resistance in Pneumonia (DRIP) score (https://www.mdcalc.com/calc/4050/drug-resistance-pneumonia-drip-score) is recommended
- 6. Recommended treatment duration for most CABP patients is 5-7 days

9. Vaccination

- 1. If the patient is not up-to-date with their vaccinations for respiratory pathogens (*Streptococcus pneumoniae*, influenza, SARS-CoV-2, RSV) they should be offered prior to hospital discharge or advised to obtain as an outpatient
 - i. Per New York Public Health Law §2805-h, vaccines for influenza and pneumococcal disease must be offered to all admitted patients age 65 or older
- 2. Providers should refer to the ACIP vaccination schedule for age appropriate and comorbidity related recommendations
 - i. https://www.cdc.gov/vaccines/schedules/hcp/index.html
- 3. Vaccines available in the hospital
 - i. S. pneumoniae (PCV20)
 - ii. Influenza
 - iii. SARS-CoV-2

<u>Table 3.</u> Antibiotic Selection for Empiric Therapy in Adults with CABP and NO Risk Factors for MRSA or *P. aeruginosa*

Non-Severe CABP Ceftriaxone 1 g IV Q24 **Azithromycin** 500 mg IV /PO Q24 **Severe CABP** Ceftriaxone 1-2 g IV Q24 **Azithromycin** 500 mg IV/PO Q24 **IV to PO Conversion for Non-Severe or Severe CABP** Amoxicillin/clavulanate 875/125 mg PO BID <u>OR</u> Cefpodoxime 200 mg PO BID + Azithromycin

500 mg PO Q24

^aFor severe beta-lactam allergy, consider levofloxacin 750 mg IV Q24

^bCan be substituted for doxycycline 100 mg IV/PO BID

^cCan be substituted for doxycycline 100 mg PO BID

<u>Table 4.</u> Antibiotic Selection for Empiric Therapy in Adults with CABP and Risk Factors for MRSA or *P. aeruginosa*

uci uginosu		
Non-Severe CABP		
Recent hospitalization & parenteral antibiotics in the last 90 days ^a	Prior respiratory isolation of MRSA ^b <u>OR</u> MRSA nasal PCR positive ^b	Prior respiratory isolation of <i>P. aeruginosa</i>
Ceftriaxone 1-2 g IV Q24 + Azithromycine 500 mg IV/PO Q24 x 3 days	Ceftriaxone 1-2 g IV Q24 + Azithromycine 500 mg IV/PO Q24 x 3 days + Vancomycin See SBUH empiric dosing guidelines	Cefepime 2 g IV Q8 <u>OR</u> Piperacillin/tazobactam 4.5 g IV Q8 + Azithromycin ^e 500 mg IV/PO Q24 x 3 days
Severe CABP		
Recent hospitalization and parenteral antibiotics in the last 90 days ^c	Prior respiratory isolation of MRSA ^b <u>OR</u> MRSA nasal PCR positive ^b	Prior respiratory isolation of <i>P.</i> aeruginosa ^d
Cefepime 2 g IV Q8 OR Piperacillin/tazobactam 4.5 g IV Q8 + Azithromycine 500 mg IV/PO Q24 x 3 days + Vancomycin See SBUH empiric dosing guidelines	Ceftriaxone 1-2 g IV Q24 + Azithromycine 500 mg IV/PO Q24 x 3 days + Vancomycin See SBUH empiric dosing guidelines	Cefepime 2 g IV Q8 <u>OR</u> Piperacillin/tazobactam 4.5 g IV Q8 + Azithromycine 500 mg IV/PO Q24 x 3 days

^aFor severe beta-lactam allergy, consider levofloxacin 750 mg IV Q24

meropenem 1 g IV Q8 + vancomycin IV

^bFor severe beta-lactam allergy, consider **levofloxacin 750 mg IV Q24 + vancomycin IV**

^cFor severe beta-lactam allergy, consider levofloxacin 750 mg IV Q24 + aztreonam 2 g IV Q8 OR

^dFor severe beta-lactam allergy, consider **levofloxacin 750 mg IV Q24 + aztreonam 2 g IV Q8** OR meropenem 1 g IV Q8

^eCan be substituted for doxycycline 100 mg IV/PO BID

Table 5. IV to PO Conversion for MRSA + P. aeruginosa

IV to PO for MRSA

Linezolid

600 mg PO BID

IV to PO for P. aeruginosa

Levofloxacin

750 mg PO Q24

IV to PO for MRSA + P. aeruginosa

Linezolid

600 mg PO BID

+

Levofloxacin

750 mg PO Q24

10. References

- Regunath H, Oba Y. Community-Acquired Pneumonia. [Updated 2020 Aug 10]. In: StatPearls [Internet]. Treasure Island (FL): StatPearls Publishing; 2020 Jan-. Available from: https://www.ncbi.nlm.nih.gov/books/NBK430749/.
- 2. Metlay JP, Waterer GW, Long AC, et al. Diagnosis and Treatment of Adults with Community-acquired Pneumonia. An Official Clinical Practice Guideline of the American Thoracic Society and Infectious Diseases Society of America. American Journal of Respiratory and Critical Care Medicine. 2019;200(7). doi:10.1164/rccm.201908-1581st.