

# Antimicrobial Stewardship Matters

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## Hospitalized CAP and HAP:

Assessment of MRSA and *P. aeruginosa* Risk, Empiric Therapy, and Duration of Therapy

### At a Glance:

#### The bottom line for hospitalized patients



1. Unless risk factors are present (see Table 1), MRSA and *P. aeruginosa* coverage is not routinely required for CAP or HAP.
2. Anti-MRSA therapy can be safely discontinued if MRSA screening is negative within 24 hours of starting anti-MRSA therapy (negative predictive value 98%)<sup>1</sup>. MRSA screening is only reliable if performed within 24 hours of initiating anti-MRSA therapy.
3. CAP therapy is typically 3 to 5 days<sup>2</sup>. HAP therapy is usually 7 days<sup>3</sup>.

### Pneumonia diagnosis

- Clinically compatible syndrome<sup>2</sup> + infiltrate on chest imaging
- CAP: acquired outside of the hospital or within 48 hours of admission
- HAP: acquired  $\geq 48$  hours after admission

### Consider transition to oral therapy when<sup>4,5</sup>

- ✓ Hemodynamically stable
- ✓ Patient is clinically improving
- ✓ Can tolerate oral intake
- ✓ Improvement in fever, respiratory status, WBC

Editor & contributors: Madison Hughes (Red Deer Regional Hospital ASP Pharmacist) and Dr. Daniel Doyle (Central Zone ASP Medical Lead).

Reviewed: Provincial AHS Antimicrobial Stewardship Committee

You can reach us at: [ahs.antimicrobialstewardship@ahs.ca](mailto:ahs.antimicrobialstewardship@ahs.ca)

Table 1: MRSA and *P. aeruginosa* Risk Factors

|                      |   |
|----------------------|---|
| MRSA                 | <ol style="list-style-type: none"> <li>1. Prior isolation of MRSA<sup>2,3</sup></li> <li>2. Broad spectrum antibiotics within the last 90 days<sup>2,3</sup></li> <li>3. MRSA consistent clinical presentation (necrotizing or cavitary pneumonia)</li> <li>4. High risk of mortality (need for ventilatory support or septic shock)<sup>3</sup></li> <li>5. Hospitalization on unit with &gt;20% of <i>S. aureus</i> isolates methicillin-resistant<sup>3</sup></li> </ol> |
| <i>P. aeruginosa</i> | <ol style="list-style-type: none"> <li>1. Prior isolation of <i>P. aeruginosa</i>, especially if from respiratory tract<sup>2,3</sup></li> <li>2. Broad spectrum antibiotics within the last 90 days<sup>2,3</sup></li> <li>3. Structural lung disease (bronchiectasis, CF)<sup>3</sup></li> </ol>  |

## Duration

**Duration:** Duration needs to be determined based on clinical response to therapy<sup>2</sup>

**CAP:** 3-5 days<sup>2,6,7</sup>      **HAP:** 7 days<sup>3</sup>

1. Three days for CAP adequate in select patients<sup>6,7</sup> (PO intake, afebrile, HR<100 bpm, RR<24 breaths/min, O<sub>2</sub> Sat≥90% on room air (or at baseline), SBP≥90 mm Hg<sup>8</sup>)
2. Longer durations indicated in severe cases<sup>2</sup> and for complicated infections (empyema, lung

Table 2: Empiric Antibiotics for Hospitalized CAP

|                            | Drug A                   | Drug B (Atypicals) | Drug C (MRSA) |
|----------------------------|--------------------------|--------------------|---------------|
| CAP                        | Ceftriaxone              | Azithromycin       |               |
| CAP + <i>P. aeruginosa</i> | Piperacillin-tazobactam* |                    |               |
| CAP + MRSA                 | Ceftriaxone              |                    | Vancomycin**  |

\*Anti-pseudomonal dosing: piperacillin-tazobactam 4.5g IV q6h, adjusted for renal function

\*\*If MRSA screening is negative, anti-MRSA therapy can be safely discontinued



Table 3: Empiric Antibiotics for HAP

|                              | Drug A  | Drug B (MRSA) |
|------------------------------|---|---------------|
| HAP ≤ 4 days hospitalization | Refer to Table 2  |               |
| HAP ≥ 4 days hospitalization | Piperacillin-tazobactam*<br>(Ceftriaxone an option if non-ICU and no <i>P. aeruginosa</i> risk factors) |               |
| HAP + MRSA                   | See above   | Vancomycin**  |

\*Anti-pseudomonal dosing: piperacillin-tazobactam 4.5g IV q6h, adjusted for renal function

\*\*If MRSA screening is negative, anti-MRSA therapy can be safely discontinued

Abbreviations: MRSA = methicillin-resistant *S. aureus*; CAP = community acquired pneumonia; HAP = hospital acquired pneumonia; WBC = white blood cells; CF = cystic fibrosis; PO = oral; HR = heart rate; RR = respiratory rate; O<sub>2</sub> Sat = oxygen saturation; SBP = systolic blood pressure; IV = intravenous

## References

<sup>1</sup>Parente DM, Cunha CB, Mylonakis E, Timbrook TT. The Clinical Utility of Methicillin-Resistant *Staphylococcus aureus* (MRSA) Nasal Screening to Rule Out MRSA Pneumonia: A Diagnostic Meta-analysis With Antimicrobial Stewardship Implications. *Clin Infect Dis*. 2018;67(1):1-7. doi:10.1093/cid/ciy024

<sup>2</sup>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. *Am J Respir Crit Care Med*. 2019;200(7):e45-e67. doi:10.1164/rccm.201908-1581ST

<sup>3</sup>Kalil AC, Metersky ML, Klompas M, et al. Management of Adults With Hospital-acquired and Ventilator-associated Pneumonia: 2016 Clinical Practice Guidelines by the Infectious Diseases Society of America and the American Thoracic Society [published correction appears in *Clin Infect Dis*. 2017 May 1;64(9):1298. doi: 10.1093/cid/ciw799] [published correction appears in *Clin Infect Dis*. 2017 Oct 15;65(8):1435. doi: 10.1093/cid/cix587] [published correction appears in *Clin Infect Dis*. 2017 Nov 29;65(12):2161. doi: 10.1093/cid/cix759]. *Clin Infect Dis*. 2016;63(5):e61-e111. doi:10.1093/cid/ciw353

<sup>4</sup>Ramirez JA, Srinath L, Ahkee S, Huang A, Raff MJ. Early switch from intravenous to oral cephalosporins in the treatment of hospitalized patients with community-acquired pneumonia. *Arch Intern Med*. 1995;155(12):1273-1276.

<sup>5</sup>Ramirez JA, Vargas S, Ritter GW, et al. Early switch from intravenous to oral antibiotics and early hospital discharge: a prospective observational study of 200 consecutive patients with community-acquired pneumonia. *Arch Intern Med*. 1999;159(20):2449-2454. doi:10.1001/archinte.159.20.2449

<sup>6</sup>el Moussaoui R, de Borgie CA, van den Broek P, et al. Effectiveness of discontinuing antibiotic treatment after three days versus eight days in mild to moderate-severe community acquired pneumonia: randomised, double blind study. *BMJ*. 2006;332(7554):1355. doi:10.1136/bmj.332.7554.1355

<sup>7</sup>Dinh A, Ropers J, Duran C, et al. Discontinuing  $\beta$ -lactam treatment after 3 days for patients with community-acquired pneumonia in non-critical care wards (PTC): a double-blind, randomised, placebo-controlled, non-inferiority trial [published correction appears in *Lancet*. 2021 Jun 5;397(10290):2150. doi: 10.1016/S0140-6736(21)01157-0]. *Lancet*. 2021;397(10280):1195-1203. doi:10.1016/S0140-6736(21)00313-5

<sup>8</sup>Hayashi Y, Paterson DL. Strategies for reduction in duration of antibiotic use in hospitalized patients. *Clin Infect Dis*. 2011;52(10):1232-1240. doi:10.1093/cid/cir063



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