### Expert Committee recommendation

The Expert Committee reviewed the evidence and limited its recommendation to hospital-acquired pneumonia. It did not include antibiotics for ventilator-associated pneumonia in this section because the condition is relatively rare and the choice of empirical antibiotic treatment in national guidelines is based on local epidemiology/microbiology. The Expert Committee endorsed the inclusion on the EML and EMLc of amoxicillin + clavulanic acid, cefotaxime and ceftriaxone for first-choice therapy in hospital-acquired pneumonia. The Committee recommended the addition of piperacillin + tazobactam to the EML and EMLc for use in hospital-acquired pneumonia as one of the first-choice therapies.

### Background

Hospital-acquired pneumonia (HAP) is defined as pneumonia with onset starting more than 48 hours after admission to hospital. Patients are often exposed to different regimens of antibiotics and thus have an increased potential to acquire resistant bacteria, making antibiotic treatment more challenging. Ventilator-associated pneumonia (VAP) is defined by the development of pneumonia while a patient is on a ventilator. Typically, the risk of infection with multidrug-resistant bacteria is high because of exposure to antimicrobials and the critical care setting. Various regimens have been assessed; a particular area of uncertainty is the need for double antipseudomonal coverage in severely ill patients. The two syndromes were combined in the application because of the relative lack of data on HAP and because the guidelines consider these together.

### Summary of evidence

A 2015 Cochrane systematic review of 6 randomized controlled trials (RCTs; 1088 participants) comparing short-course with long-
Guidelines

Rationale for antibiotic selection

Amoxicillin + clavulanic acid is a core antibiotic that can be used within 5 days of hospital admission and if there is no prior antibiotic exposure or risk for resistance. Third-generation cephalosporins are another core choice, as is piperacillin + tazobactam. The systematic reviews suggest non-inferiority between vancomycin and linezolid. Linezolid, however, was not proposed as a core antibiotic since it is proposed for the preservation list of those antibiotics that are last-line for highly resistant pathogens. Use of empirical vancomycin should be restricted to cases where MRSA is suspected. Aminoglycosides are on the list for double antipseudomonal coverage if needed. The application proposed ceftazidime, cefepime and piperacillin + tazobactam for antipseudomonal coverage. It is recommended that the fluoroquinolones be used only when needed, for example, in the case of a serious allergy to first-choice antibiotics. Given the concern about carbapenem resistance, these agents should be used only when there are no other alternatives.

Committee considerations

For common community-acquired infections, the main focus has been on empirical treatment choices that are broadly applicable in most countries. Generally, alternatives for use in case of allergy were not considered. The Committee decided not to include VAP in this review because of the need to have local microbiology and epidemiological data to guide the choice of antibiotics and because it is a relatively rare condition. The Committee considered the various antibiotics proposed in the application under the guiding principle of parsimony and selected first-choice antibiotics for HAP for inclusion on the EML and/or EMLc. As a result, levofloxacin,
moxifloxacin, ciprofloxacin, ceftazidime, aztreonam, meropenem, imipenem, amikacin, gentamicin, tobramycin and vancomycin were excluded. Recommended first-choice antibiotics are reported below. The first-choice antibiotics are those generally recommended based on available evidence and are usually narrow-spectrum agents.

### EML recommendations: Other specified pneumonia

<table>
<thead>
<tr>
<th>First choice</th>
<th>Second choice</th>
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<tbody>
<tr>
<td><strong>HOSPITAL-ACQUIRED PNEUMONIA</strong></td>
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<td>Piperacillin + tazobactam</td>
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<td>Amoxicillin + clavulanic acid</td>
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<td>Cefotaxime</td>
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