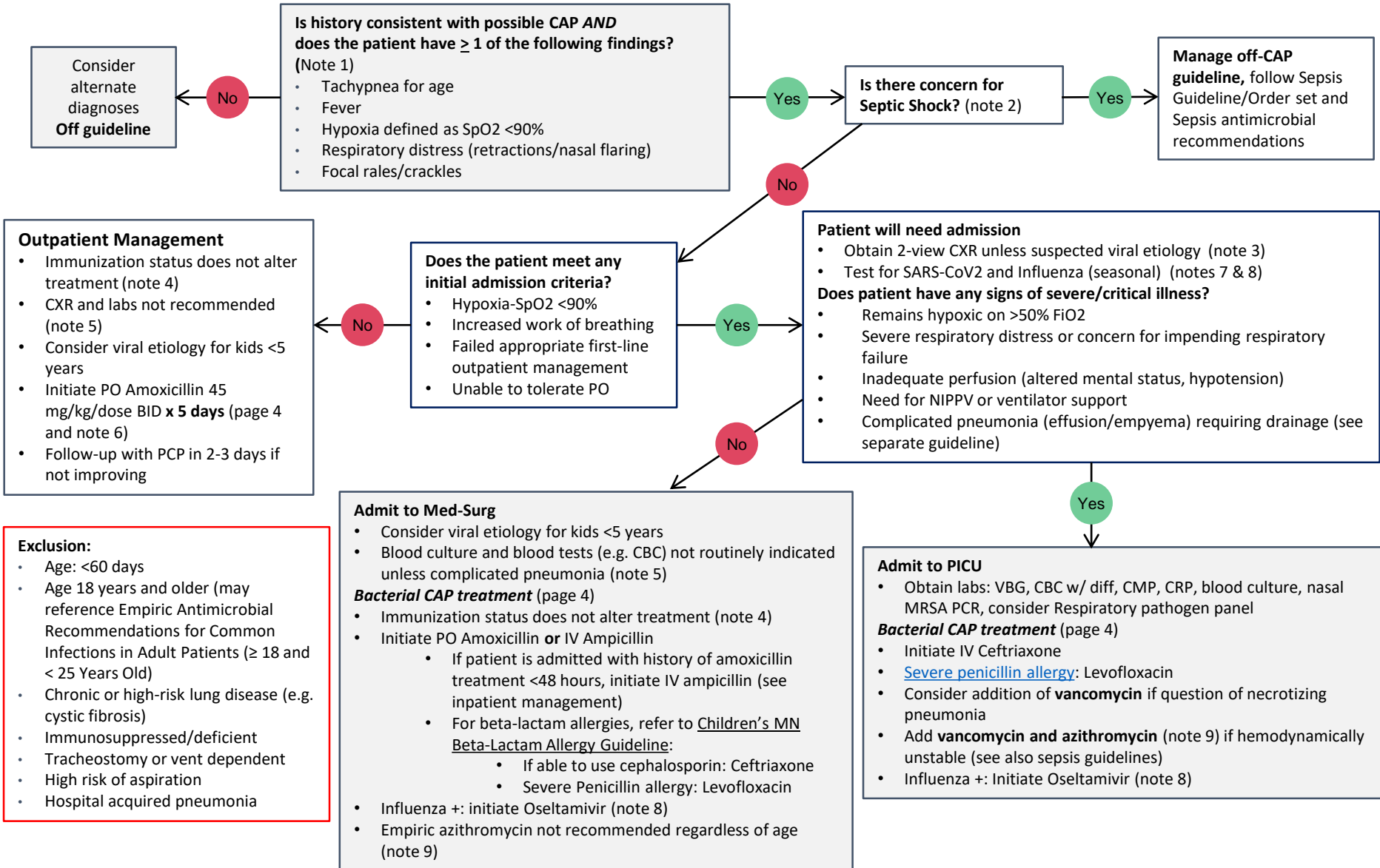
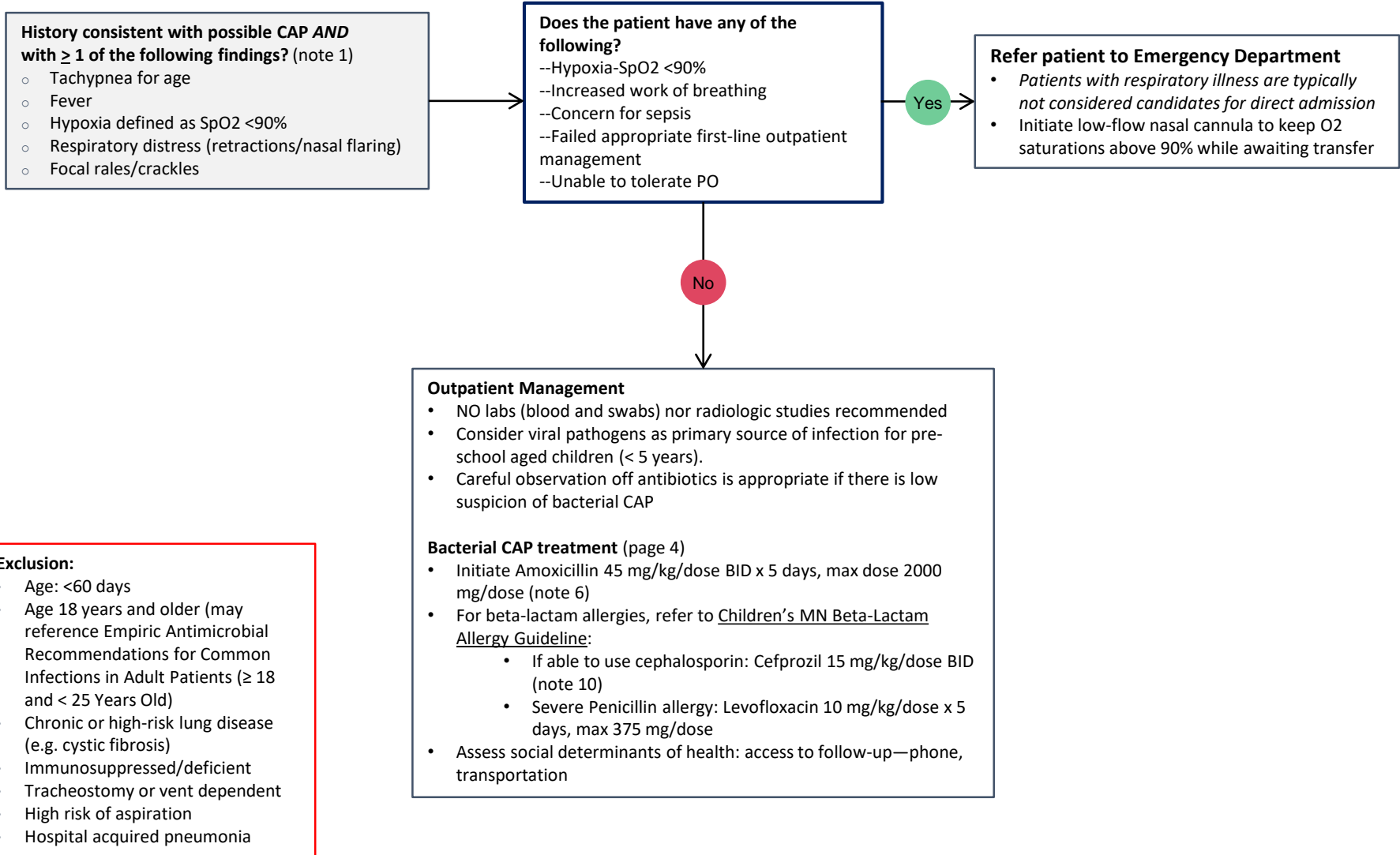


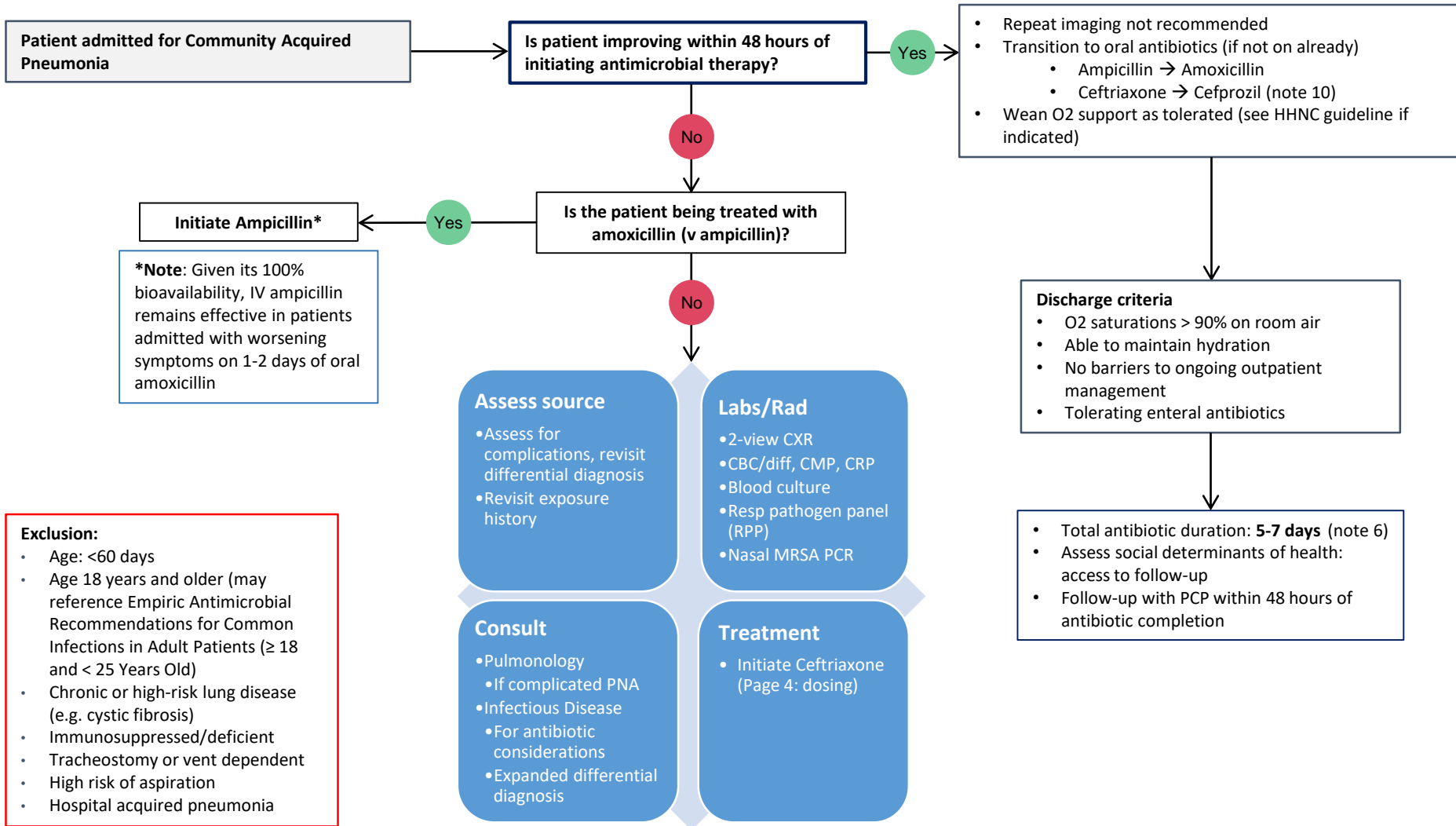
Aim: To decrease variation in the treatment of CAP.



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Exclusion:

- Age: <60 days
- Age 18 years and older (may reference Empiric Antimicrobial Recommendations for Common Infections in Adult Patients (≥ 18 and < 25 Years Old))
- Chronic or high-risk lung disease (e.g. cystic fibrosis)
- Immunosuppressed/deficient
- Tracheostomy or vent dependent
- High risk of aspiration
- Hospital acquired pneumonia

Antimicrobial	Dosing	Interval	Max single dose
Amoxicillin (PO)	45 mg/kg/dose	BID	2,000 mg
Ampicillin (IV)	50 mg/kg/dose	q6h	2,000 mg
Cefdinir (PO)	7 mg/kg/dose	BID	300 mg
Cefprozil (PO)	15 mg/kg/dose	BID	500 mg
Ceftriaxone (IV) Hemodynamically-stable	50 mg/kg/dose	q24h	2,000 mg
Hemodynamically unstable		q12h	
Levofloxacin (IV/PO) 6 mos to 4 years	10 mg/kg/dose	BID	375 mg
>4 years		Once daily	750 mg
Osetamivir (PO) <i>Red Book</i> Birth to <12 months 9-11 months 1-12 years ≤15 kg 15.1-23 kg 23.1-40 kg >40 kg ≥13 years	3 mg/kg PO 3.5 mg/kg 30 mg PO 45 mg PO 60 mg PO 75 mg PO 75 mg PO	BID	See Dosing

Beta-lactam notes:

- High-dose amoxicillin** (90 mg/kg/day) **divided BID** is expected to be adequate for the treatment of uncomplicated pneumonia in the majority of children given the low rates of *Streptococcus pneumoniae* resistance to penicillin locally. Recent US clinical trials on pediatric CAP have used BID dosing ([SCOUT-CAP](#)).
- Please see [Children's Minnesota Beta-Lactam Allergy Guideline](#) for further information:
 - Severe Penicillin Allergy** definition: includes any of the following: anaphylaxis, angioedema, cardiac arrest, respiratory distress, interstitial nephritis, serum sickness, severe cutaneous reaction (e.g. Stevens-Johnson syndrome, erythema multiforme, DRESS, and TEN.)
 - Nonsevere allergic reactions to amoxicillin**, these are the options per the AAP 2011 CAP guideline: trial of amoxicillin under medical observation; trial of cephalosporin under medical supervision; treatment with levofloxacin.

Disclaimer: This guideline is designed for general use with most patients; each clinician should use their own independent judgment to meet the needs of each individual patient. This guideline is not a substitute for professional medical advice, diagnosis or treatment.

Note 1. Community acquired pneumonia

- Typical bacterial pathogens that cause CAP include *Streptococcus pneumoniae*, *Haemophilus influenzae*, and *Moraxella catarrhalis*. However, with the advent of novel diagnostic technologies, *viral* respiratory pathogens are increasingly being identified as frequent etiologies of CAP.
- There is no universal presentation of pneumonia. Overlapping features occurring in various lower respiratory tract conditions (e.g., asthma, bronchiolitis, pneumonitis, pneumonia) makes the clinical diagnosis challenging. Common signs and symptoms include fever, cough, tachypnea, increased work of breathing, dehydration and hypoxia. Other symptoms can include abdominal pain and emesis. Physical exam findings may include rales, crackles, wheezing, dyspnea as demonstrated by retractions, nasal flaring or grunting, and decreased breath sounds.

Note 2. Sepsis: signs and symptoms

- Core temp abnormality: • T > 38.4°C • Or T < 36.0°C in any age PLUS 1 or more of the following: • High-risk condition • Change in perfusion: grey, cool, mottled, clammy or flush • Change in mental status: agitation, distress, inc: 5yonsolable, lethargic, limp • Hypotension (MAP ≤ 5th %ile for age)
- Additional signs include: • Tachycardia • Reduced urine output • Tachypnea/new O2 requirement • High caregiver concern
- Severe Sepsis:
 - Sepsis + CV dysfunction or ARDS or 2+ organ dysfunctions
 - (Also referred to as Sepsis-Associated acute organ dysfunction)
- Septic Shock: Sepsis + CV dysfunction that persists after ≥40 mL/kg NS in one hour

Note 3. CXR considerations: No need to repeat if imaging from outside facility is readily available for review or strong suspicion of viral illness.

Note 4. Immunization status. This guideline recommends that treatment utilize same practice for vaccinated and unvaccinated patients. Immunization status influenced antibiotic choices in the 2011 IDSA/PIDS CAP guideline. However, these guidelines were composed during the widespread use of the pneumococcal conjugate vaccine 7 (PCV7), introduced in 2000. The adoption of PCV13, beginning in 2010, had a further effect on clonal circulation and herd immunity. This has led to: 1. Reduction in Pneumococcal pneumonia hospital admissions in vaccinated *and* unvaccinated children; 2. Reduction in penicillin-resistant invasive pneumococcal infections; 3. rare HiB-associated pneumonias

Note 5. Blood culture: Multiple studies reviewing utility of BC in management of CAP--5-yr retrospective cohort study reviewing >7500 non-ICU hospitalized children found only a 2.5% positive rate (78% positive with Pneumococcus; 82% penicillin susceptible). Prevalence of bacteremia among severe or complicated PNA was 4.2% v 2.2%.

Note 6. Short course antibiotics for treatment of CAP: 5-7 days is recommended.

- SAFER (blinded RCT) trial reviewed short v long course (5 days v 10 days) Amoxicillin for non-hospitalized CAP had comparable clinical cure rates
- Same et al. (2020) Retrospective review of non-ICU hospitalized children comparing short v long (5-7d b v 8-14 days) antibiotic course had no difference in treatment failure between two therapy duration groups (3% v 6%)
- 5 day duration is adequate in most hospitalized patients who are improving and have reached clinical stability by day 5 of treatment. 7 day duration may be favored in hospitalized patients who are slower to respond to initial therapy, and is indicated for those with CAP due to *MRSA* or *Pseudomonas aeruginosa*.

Note 7. COVID-19 considerations: continue to follow local Children's Minnesota guidance for outpatient and inpatient testing.

Note 8. Influenza considerations: Influenza-Like Illness (ILI): Fever $\geq 100.4^{\circ}\text{F}$ and Cough and/or Sore Throat; During Influenza Season

- Starting Oseltamivir: High risk patients: ≤ 12 months old or chronic conditions such as pulmonary (e.g., asthma), cardiac, renal, hepatic, hematologic, metabolic, neurologic; Immunosuppressed; long term ASA therapy; morbid obesity; social factors; resident of chronic care facility; household contact < 6 months or with chronic condition

Note 9. Azithromycin: Empiric use of azithromycin for coverage of atypical pathogens (e.g. *Mycoplasma pneumoniae*) is **not recommended** for the majority of pediatric patients regardless of patient age.

- A large observational study showed no benefit of empiric combination therapy with a macrolide plus beta lactam compared with beta lactam monotherapy in hospitalized children (<18 years of age) with radiographically confirmed CAP with regards to length of stay, re-hospitalizations, or recovery at follow-up.⁹ Lack of benefit of macrolide therapy was also shown among a subgroup of hospitalized children with atypical bacteria detected

Note 10. Cephalosporin availability and insurer coverage: Historically, insurer coverage for 2nd/3rd generation cephalosporins is variable—which places families at risk for out-of-pocket (OOP) cost.

- Cefprozil is usually covered by insurance. Children's Minnesota Outpatient Pharmacy stocks 250mg/5ml suspension and 500 mg tablets.
- Alternatively, if Cefprozil is not ideal for OOP cost and/or availability, Cefdnir may be used.

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Clinical Guidelines: Seattle Children's and Children's Hospital of Philadelphia

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