COMMUNITY ACQUIRED PNEUMONIA (CAP)

Intended for patients age >60 days to <18 years with signs/symptom suggestive of pneumonia (guidance can be extrapolated down to a healthy term newborn)

Exclusion Criteria:

- Immunocompromised
- Tracheostomy
- Chronic conditions including:
- Cystic fibrosis
- •BPD on baseline supplemental oxygen
- Sickle cell disease
- Neuromuscular disease with baseline respiratory compromise
- Unrepaired cyanotic heart disease
- Large/clinically significant^A effusion/empyema/lung abscess

Signs/Symptoms of CAP B

(*highest predictive value)

- Fever >101° F
- Tachypnea for age*
 - 2-12 months: RR>50
 - 1-5 years: RR>40
 - 5 years-adolescents: RR>25
 - Adolescents: RR>20
- SpO2 <90%*
- Increased work of breathing: nasal flaring*, retractions, belly breathing, tracheal tugging, grunting, head bobbing
- Breath sound abnormalities: focal findings
- Chest pain
- Abdominal pain
- Preceding upper respiratory infection
- Symptoms cannot be explained by bronchiolitis (Signs/symptoms of bronchiolitis include: age <2 years, variable and dynamic disease process, wheezing, upper respiratory infection followed by lower respiratory signs with peak on day 3-5)

Assess severity of symptoms

Mild Criteria (Outpatient criteria)

All of the following:

- Non-toxic appearing
- O2 sats ≥90% on room air
- Tolerating work of breathing^C
- Able to maintain adequate hydration
- Able to tolerate enteral medications

See outpatient algorithm

Moderate Criteria (Acute care hospitalization criteria)

One or more of the following AND does not meet severe/critical care criteria:

- Need for supplemental O2 to
- maintain saturations ≥90%
 Not tolerating work of breathing^c
- Need for IV fluids to maintain adequate hydration
- Inability to tolerate enteral meds

See hospital algorithm

Severe Criteria (Critical Care Criteria)

One or more of the following:

- Invasive mechanical ventilation
- CPAP/BiPAP requirement
- Oxygen requirement >10L oxymask

See hospital algorithm

Exit pathway, contact Critical Care as needed

For questions concerning this work, contact cappathway@childrenswi.org

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Return to: (main algorithm)

Original: 11/2024 Revised: 4/2025

Revised: 4/2025
Process owner: CAP Pathway Team

COMMUNITY ACQUIRED PNEUMONIA (CAP)

Outpatient Pathway

(including CMG/Outpatient/Urgent Care/ED)

Intended for patients age >60 days to <18 years with signs/symptom suggestive of pneumonia (guidance can be extrapolated down to a healthy term newborn)

Mild criteria (outpatient/urgent care/ED)

All of the following:

- Non-toxic appearing
- O2 sats ≥90% on room air
- Tolerating work of breathing^C
- Able to maintain adequate hydration
- Able to tolerate enteral medications



- **Diagnostic testing** Imaging: routine imaging not recommended
- Serum studies: routine CBC/inflammatory markers, blood culture not recommended
- Viral swabs: consider influenza/COVID 19 testing seasonally/as appropriate (Information for Health Professionals | CDC). Additional testing/Respiratory Pathogen Panel not generally recommended



Treatment

- Amoxicillin 90 mg/kg/day divided TID, max 3 g daily
- 5 days duration if able to be reassessed in 3-4 days
- Consider 7 days if uncertain follow-up in 3-4 days
- See table for additional antibiotic considerations including allergy recommendations



Follow Up

- EDTC/UC: Patient to call PCP or specialist next business day to discuss follow-up plans
- CMG/Ambulatory: Follow up in 3-4 days; sooner if worsening. Follow up may be in-person, via phone or MyChart, or prn if not improving, based on provider's clinical judgment
- Patient may return to school or child care when afebrile for 24 hours, adequately hydrated, and tolerating other symptoms. Cough may persist and does not require exclusion from school or child care
- Repeat imaging rarely needed. Consider if extenuating circumstances (e.g. concern for scarring). If obtained, wait at least 4 weeks prior to repeating

Exclusion Criteria:

- Immunocompromised
- Tracheostomy
- Chronic conditions including:
 - Cystic fibrosis

No

- •BPD on baseline supplemental oxygen
- •Sickle cell disease
- Neuromuscular disease with baseline respiratory compromise
- •Unrepaired cyanotic heart disease
- Large/clinically significant^A effusion/empyema/lung abscess

Moderate criteria (hospitalized/ED)

>/= 1 of the following AND does not meet severe critical care criteria:

- Need for supplemental O2 to maintain saturations ≥90%
- Not tolerating work of breathing^C
- Need for IV fluids to maintain adequate hydration
- Inability to tolerate enteral medications



Consider hospitalization

- For questions regarding management or hospitalization at the Milwaukee campus → call Physician Consult Center (414-266-2470)
- For questions regarding management or hospitalization in Fox Valley→page the FV Peds Hospitalist team (920-554-
- See considerations for direct admission to acute care hospitalization

Treatment failure recommendations (lack of clinical improvement after 72 hours of

appropriate antimicrobial therapy)E Do not hospitalize solely for treatment failure if not also

- meeting moderate criteria for hospitalization ■ If true treatment failure AND meeting moderate criteria for
- hospitalization, start IV ampicillin. Do not need broader coverage based solely on outpatient treatment failure regardless of what enteral antibiotic was used as IV route may be sufficient escalation of treatment.
- Consider possibility of viral pneumonia and reconsider need for ongoing antibiotics
- Ensure correct antibiotic and dose/frequency of antibiotic
- Ensure patient is taking antibiotic as prescribed
- In times of high community prevalence, consider alternative diagnoses such as pertussis or atypical pneumonia
- If persistent fevers, consider CXR to confirm diagnosis and assess for clinically significant effusion^A
- <u>Do not add azithromycin</u>F or broaden coverage including amoxicillin/clavulanic acid or cephalosporin based solely on concern for treatment failure
- If questions regarding need for alternative antimicrobial management, consider contacting Antimicrobial Stewardship provider via EPIC Secure Chat (preferred) or page. For holidays/after-hours, page on-call ID provider^E
- If questions regarding need for airway clearance or alternative diagnosis, contact on call pulmonary provider

Not improving

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COMMUNITY ACQUIRED PNEUMONIA (CAP)

Hospitalized/ED Pathway

Intended for patients age >60 days to <18 years with signs/symptom suggestive of pneumonia (guidance can be extrapolated down to a healthy term newborn)

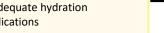
Exclusion Criteria:

- Immunocompromised
- Tracheostomy
- Chronic conditions including:
- Cystic fibrosis
- •BPD on baseline supplemental oxygen
- •Sickle cell disease
- Neuromuscular disease with baseline respiratory compromise
- •Unrepaired cyanotic heart disease
- Large/clinically significant^A effusion/empyema/lung abscess

Moderate criteria (acute care hospitalization/ED)

One or more of following AND does not meet severe critical care criteria:

- Need for supplemental O2 to maintain saturations ≥90%
- Not tolerating work of breathing^c
- Need for IV fluids to maintain adequate hydration
- Inability to tolerate enteral medications



No

Severe criteria (critical care hospitalization/ED)

Diagnostic testing

· Serum studies: recommend CBC and blood culture; consider

appropriate (Information for Health Professionals | CDC)

Viral swabs: consider influenza/COVID 19 testing seasonally/as

MRSA nasal PCR if starting MRSA coverage (see treatment below)

One or more of the following:

 Invasive mechanical ventilation ■ CPAP/BiPAP requirement

Imaging: CXR (PA and lateral)

ESR/CRP/procalcitonin

Oxygen requirement >10L oxymask

Diagnostic testing

- Imaging: CXR (PA and lateral)
- Serum studies: routine CBC/inflammatory markers, blood culture not recommended
- Viral swabs: consider influenza/COVID 19 testing seasonally/as appropriate (Information for Health Professionals | CDC). Additional testing/Respiratory Pathogen Panel not generally recommended

Treatment

- Ampicillin 50 mg/kg/dose q6h, max 8g daily
- Bronchial hygiene protocol^G
- See table for additional antibiotic considerations



Improving

Discharge criteria

- Non-toxic appearing
- O2 sats ≥ 90% on RA for at least 6 hours
- Improving fever curve
- Tolerating work of breathing^c
- Able to maintain adequate hydration and tolerate oral/enteral medications



Treatment

- Hemodynamically stable: Ceftriaxone +/- vancomycin or clindamycin based on clinical judgement. If MRSA coverage is started, de-escalate if either MRSA nasal PCR negative or ET tube
- culture negative for MRSA • Hemodynamic instability: Ceftriaxone + vancomycin. De-escalate
- vancomycin when blood culture negative for MRSA and either MRSA nasal PCR negative or ET tube culture negative for MRSA
- Bronchial hygiene protocol^G
- See table for additional antibiotic considerations

Discharge recommendations

- Amoxicillin 90 mg/kg/day divided TID, max 3 g daily
- Consider 7-day total antibiotic course for moderate-severe cases
- 5 days may be appropriate if mild/rapid improvement
- Follow up in 2-3 days

Original: 11/2024

- Patient may return to school or child care when afebrile for 24h, adequately hydrated, and tolerating other symptoms. Cough may persist and does not require exclusion from school or child care
- Repeat imaging rarely needed. Consider if extenuating circumstances (e.g. concern for scarring). If obtained, wait at least 4 weeks prior to repeating

<u>Treatment failure recommendations</u>^E

Failing ampicillin inpatient:

- Lack of clinical improvement at 48hrs: Obtain CRP or procalcitonin (if not already obtained). Consider obtaining Respiratory Pathogen Panel and continue current therapy
- Lack of clinical improvement at 72hrs: Obtain repeat CXR and repeat CRP or procalcitonin; if no clinical/lab improvement, consider escalation in therapy (ie: ceftriaxone)); if clinically significant effusion^A -> off Pathway, see CAP with PE guideline; if Respiratory Pathogen Panel positive for mycoplasma pneumonia, consider treatment with azithromycin^F

For questions concerning this work, contact

Revised: 4/2025 Process owner: CAP Pathway Team

Antimicrobial considerations		
Aspiration Concern	See CW aspiration pneumonitis vs pneumonia guideline for guidance if concern for aspiration pneumonia	
Atypical pneumonia	• The use/addition of azithromycin is not routinely recommended for CAPF. Could consider if concern for atypical etiologies (>4 years old, slowly progressive symptoms, extrapulmonary findings including rash, GI symptoms). Atypical etiologies are unlikely in patients under 5 years. B	
Ceftriaxone transition to enteral option	 If started for severe CAP/PICU→ most likely etiology is strep pneumonia. Can likely transition to amoxicillin. Consider amoxicillin/clavulanate if slow to improve and/or based on clinical judgement If started for treatment failure after 72 hours of ampicillin→ amoxicillin/clavulanate If started for low-risk penicillin allergy^H→ 1st line: cefprozil or 2nd line: cefuroxime If started for high-risk penicillin allergy^H→ 1st line: cefuroxime or 2nd line: clindamycin Local ID experts do not recommend use of cefdinir for the treatment of CAP^I 	
Co-infections	 Acute otitis media: Concurrent acute otitis media may impact the recommended duration of treatment for pneumonia. Current AAP guidelines⁷ for AOM recommend 10-day course for children younger than 2 years and all children with severe symptoms, 7-day course is recommended in children 2 years of age and older with mild or moderate AOM. If treating for concurrent acute otitis media and CAP, frequency should be TID dosing Conjunctivitis:	
Immunization status	 No changes in diagnostics or treatment are indicated based solely on immunization status including pneumococcal status. Following the implementation of pneumococcal vaccination, invasive pneumococcal disease and complicated pneumococcal pneumonia have decreased substantially for al children regardless of vaccination status. Complicated pneumonia is more likely to be due to non- vaccine related strains. In addition, local sensitivity rates for strep pneumonia are very high (<u>Children's Wisconsin Antibiogram</u>). Local ID experts therefore do not recommend change in management based solely on immunization status. 	
Influenza positive	 Mild/moderate pneumonia Although there is an association between influenza and subsequent staph aureus pneumonia, the overall risk is still lower than strep pneumonia. Addition of clindamycin is not needed in mild/moderate pneumonia solely based on recent influenza infection unless other clinical concerns. Severe pneumonia: If severe pneumonia and influenza positive, local ID experts recommend ceftriaxone + vancomycin OR clindamycin. Can de-escalate vancomycin/clindamycin MRSA coverage when either MRSA nasal PCR negative or ET tube culture negative for MRSA. 	
MRSA coverage	If antimicrobial coverage is escalated to include MRSA coverage (vancomycin/clindamycin), obtain MRSA nasal PCR for de-escalation if negative.	
	See next page for additional antimicrobial considerations	

See next page for additional antimicrobial considerations

Original: 11/2024 Revised: 4/2025 For questions concerning this work, contact

Antimicrobial considerations (Continued)

Penicillin allergies

Low-risk penicillin allergy^H:

- For hospitalized patients, if low-risk penicillin questionnaire (located in admission tab), can give ampicillin if family agreeable to penicillin challenge. The patient would then need to do the one dose oral trial of the medication they previously reacted to prior to discharge. Contact Antimicrobial stewardship team in EPIC Secure Chat if questions on penicillin oral challenge protocol.
- For outpatient, can refer to the Penicillin Delabeling Clinic as applicable for future illnesses
- IV: ceftriaxone or ampicillin if family agreeable to penicillin challenge
- Enteral:
 - 1st line: cefprozil (suspension or tablets)
 - 2nd line: cefuroxime (tablets, suspension not available) Tablets may be crushed and added to food, but can cause bitter taste
 - 3rd line: clindamycin (liquid or capsules) if above not available. Liquid clindamycin can be poorly tolerated due to taste. Consider prescribing capsules which can be opened and mixed in small amount of food to increase adherence
 - See enteral antibiotic table for additional specifics on cephalosporin alternatives
 - Local ID experts do not recommend use of cefdinir for the treatment of CAP^I

High-risk penicillin allergy^H:

- IV: ceftriaxone
- Enteral
 - 1st line: cefuroxime tablets (suspension not available). Tablets may be crushed and added to food, but can cause bitter taste
 - 2nd line: clindamycin (liquid or capsules). Liquid clindamycin can be poorly tolerated due to taste. Consider prescribing capsules which can be opened and mixed in small amount of food to increase adherence
 - See enteral antibiotic table for additional specifics on cephalosporin alternatives
 - Local ID experts do not recommend use of cefdinir for the treatment of CAPI

Treatment Failure^E: (lack of

improvement

after 72h of

appropriate antimicrobial

therapy)

clinical

Outpatient amoxicillin treatment failure:

- Do not hospitalize solely for treatment failure if not also meeting moderate criteria for hospitalization
 If true treatment failure AND meeting moderate criteria for hospitalization, start IV ampicillin. Do not need broader coverage based solely on outpatient treatment failure regardless of what enteral antibiotic was used as IV route may be sufficient escalation of treatment
- Consider possibility of viral pneumonia and reconsider need for ongoing antibiotics
- Ensure using correct antibiotic and dose/frequency of antibiotic
- Ensure patient is taking antibiotic as prescribed
- In times of high community prevalence, consider diagnoses such as pertussis or atypical pneumonia
- If persistent fevers, consider CXR to confirm diagnosis and assess for clinically significant effusion
- <u>Do not add azithromycin^F</u> or broaden coverage including amoxicillin/clavulanic acid or cephalosporin based solely on concern for treatment failure
- If questions regarding need for alternative antimicrobial management, consider contacting Antimicrobial Stewardship provider via EPIC Secure Chat (preferred) or page. For holidays/after-hours,
- page on-call ID provider^E
 If questions regarding need for airway clearance or alternative diagnosis, contact on-call pulmonary

Hospitalized ampicillin treatment failure:

- Lack of clinical improvement at 48h: Obtain CRP or procalcitonin (if not already obtained), consider obtaining Respiratory Pathogen Panel and continue current therapy
- Lack of clinical improvement at 72hrs: Obtain repeat CXR and repeat CRP or procalcitonin; if no clinical/lab improvement, consider escalation in therapy (ie: ceftriaxone); if clinically significant effusion^A -> off Pathway, see CAP with PE guideline; if Respiratory Pathogen Panel positive for mycoplasma pneumonia, consider treatment with azithromycin^F
- If questions regarding need for alternative antimicrobial management, contact Antimicrobial Stewardship provider^E
- · If questions regarding need for airway clearance or alternative diagnosis, contact on call pulmonary

For questions concerning this work, contact

Original: 11/2024 Revised: 4/2025

clindamycin, and 40% of azithromycin in the Childrens Wisconsin system.

Parenteral antimicrobial dosing				
Medication	Dose	Max dose		
Ampicillin	50 mg/kg/dose q6	Max 8 grams daily		
Ceftriaxone	100 mg/kg/dose q24	Max 2 grams daily		
Clindamycin	10 mg/kg/dose q6	Max 2.7 grams daily		
Vancomycin	Pharmacy to dose; see Vancomycin Cli	inical Practice Guideline for more info		

Medication	Dose	Max dose	Other
Wicalcation		First Line*	Other
A t - 1111	OO marker (days divided TID		dditional information on formulations and may desire
Amoxicillin	90 mg/kg/day divided TID	*TID dosing optimizes time of k improve compliance and may b instances: 90mg/kg/day divided	 idditional information on formulations and max dosing iilling and is generally preferred. BID dosing may be necessary in some social situations. In those d BID
		Alternatives*	
Amoxicillin/	90 mg/kg/day divided TID	See amoxicillin/clavulanate guideline for additional information on	
clavulanate (14:1)		formulations and max dosing	
Cephalosporins	Cefprozil 30mg/kg/day divided BID	Max 1000mg daily	Local ID experts do not recommend use of cefdin for the treatment of CAP ^I
			Cefprozil is covered by Medicaid, is typically mos cost-effective, and has better pharmacokinetic than other cephalosporins so is preferred if
	Cefuroxime 30mg/kg/day divided BID (only available in tablet form)	Max 1000mg daily	available. It shares a side chain with amoxicillin s is not appropriate alternative for high-risk penicillin allergy .
			Cefuroxime is an appropriate alternative if cefprozil is not available and for high-risk penicillin allergy! patients. It is only available in tablet form. Tablets can be crushed/mixed thoug can cause bitter taste. Ensure pharmacy has in stock and medication is not cost prohibitive.
			If issues finding cefprozil, see pharmacy consideration table.
Clindamycin	30 mg/kg/day divided q8	Max 1800mg daily	Liquid clindamycin can be poorly tolerated due to taste. Consider prescribing capsules which can be opened and mixed in small amount of food to increase adherence

EHR Tools			
Admission order set	"Gen peds pneumonia admission"		
Discharge order set/AVS template	"Gen DC pneumonia"		
Amoxicillin and Augmentin Dosing for Discharge Order set	"Amoxicillin and Augmentin High and Standard Dosing for Discharge" within discharge order sets		
Outpatient antibiotic order panel	Community acquired pneumonia antibiotics smart order panel (search "pneumonia"). UC version is noted by "(urgent care only)"		
Outpatient AVS (can start with .avs, .ucdc, or .pcav)	.avspneumoniabacterial, .avspneumoniaviral, .avspneumoniaatypical; Spanish versions end in SP		
CMG/Urgent Care sick visit plan block	Find in note template plan list as "pneumonia" or use smartphrase planpneumonia		

For questions concerning this work, contact

Return to: (main algorithm)

Original: 11/2024

Revised: 4/2025

Pharmacy Considerations

Skywalk Pharmacy frequently carries the enteral medications recommended in this pathway. Skywalk Pharmacy is open Monday-Friday 8am-11pm and Saturday/Sunday/holidays 8am-6pm.

Antimicrobial Stewardship is partnering with Walgreens to support stocking cefprozil suspension/ tablets at the following locations.

CW Main Campus	Nearest pharmacy Walgreens	
Childrens Wisconsin Main Campus	Skywalk Pharmacy: 8915 W. Connell Ct. Milwaukee, WI 53226	
8915 W. Connell Ct., Milwaukee, WI 53226	Walgreens: 2275 N Mayfair Rd, Wauwatosa, WI 53226	
CW Urgent Care	Nearest Walgreens	
Kenosha UC	7520 118th Ave, Pleasant Prairie, WI 53158	
6809 122 Ave, Kenosha, WI 53142		
Forest Home UC	1433 W Burnham St, Milwaukee, WI 53204	
1432 W Forest Home Ave, Milwaukee, WI 53204		
Mayfair UC	2275 N Mayfair Rd, Wauwatosa, WI 53226	
3040 N 117th St #200, Wauwatosa, WI 53222		
Delafield UC	2901 Golf Rd, Delafield, WI 53018	
3195 Hillside Dr, Delafield, WI 53018		
Mequon UC	11270 N Port Washington Rd, Mequon, WI 53092	
1655 W Mequon Rd, Mequon, WI 53092		
New Berlin UC	3855 S Moorland Rd, New Berlin, WI 53151	
4855 S Moorland Rd, New Berlin, WI 53151		
CW Primary Care	Nearest Walgreens	
Bayshore Pediatrics (Fox Point)	8615 N Port Washington Rd, Fox Point, WI 53217	
7950 N Port Washington Rd, Fox Point, WI 53217		
Cedarburg Pediatrics	W62n190 Washington Ave, Cedarburg, WI 53012	
7861 WI-60 Trunk, Cedarburg, WI 53012	1915 Wisconsin Ave, Grafton, WI 53024	
North Shore Pediatrics (Mequon)		
1655 W Mequon Rd, Mequon, WI 53092		
River Glen Pediatrics (Glendale)	5400 N Port Washington Rd, Glendale, WI 53217	
4655 N Port Washington Rd, Glendale, WI 53212	320 E Capitol Dr, Milwaukee, WI 53212	
West Bend Pediatrics	1720 W Washington St, West Bend, WI 53095	
611 Veterans Ave #106, West Bend, WI 53090		
Forest Home Pediatrics		
Forest View Pediatrics (New Berlin)	3855 S Moorland Rd, New Berlin, WI 53151	
4855 S Moorland Rd, New Berlin, WI 53151		
Franklin Pediatrics	7130 S 76th St, Franklin, WI 53132	
7322 W Rawson Ave, Franklin, WI 53132		
Kenosha Pediatrics		
Lakeside Pediatrics (Kenosha)	7520 118th Ave, Pleasant Prairie, WI 53158	
8600 75th St, Kenosha, WI 53142	7535 Green Bay Rd, Kenosha, WI 53142	
Oak Creek Pediatrics	9449 S Howell Ave, Oak Creek, WI 53154	
8375 S Howell Ave, Oak Creek, WI 53154		
Greenfield Pediatrics	2677 S 108th St, West Allis, WI 53227	
3365 S 103rd St, Milwaukee, WI 53227	9100 Beloit Rd, Milwaukee, WI 53227	
Southwest Pediatrics (New Berlin)		
Good Hope Pediatrics	9040 Good Hope Rd, Milwaukee, WI 53224	
7720 Good Hope Rd, Milwaukee, WI 53223	6442 N 76th St, Milwaukee, WI 53223	
Mayfair Pediatrics (Wauwatosa)	2275 N Mayfair Rd, Wauwatosa, WI 53226	
3040 N 117th St, Wauwatosa, WI 53222	10800 W Capitol Dr, Wauwatosa, WI 53222	
Midtown Pediatrics	5115 W Capitol Dr, Milwaukee, WI 53216	
5433 W Fond du Lac Ave, Milwaukee, WI 53216		
Bluemound Pediatrics (Brookfield)	15350 W Bluemound Rd, Elm Grove, WI 53122	
12635 W Bluemound Rd, Brookfield, WI 53005		
Delafield Pediatrics		
Pewaukee Pediatrics	1441 Capitol Dr, Pewaukee, WI 53072	
1215 George Towne Dr, Pewaukee, WI 53072		
Westbrook Pediatrics (Brookfield)	15640 W Capitol Dr, Brookfield, WI 53005	
14755 W Capitol Dr #100, Brookfield, WI 53005		

For questions concerning this work, contact

Appendix A: Large/clinically significant^ effusion/empyema/lung abscess

- Large or clinically significant effusion (excluded from pathway managed on CAP with PE guideline):
 - >10 mm on decubitus x-ray or chest ultrasound or greater than ¼ of hemithorax opacified AND requiring drainage or chest tube placement after discussion with Interventional Radiology (IR)
- •Clinically insignificant effusion (included in pathway-no change in management from patients without effusion):
 - •<10 mm on decubitus x-ray or chest ultrasound or less than ¼ of the hemithorax opacified AND not requiring drainage or chest tube placement*
 - *if question about the need for drainage, recommend discussion with IR
- Instructions for contacting IR:
 - •Obtain 2 view AP/lateral chest x-ray. Decubitus imaging not required
 - Contact IR regarding chest tube placement and if additional imaging (US, CT) indicated prior to placement
 - •IR may also choose to obtain US as part of chest tube placement. Additional US order not needed in those cases

Appendix B: Differentiating bacterial vs viral vs atypical pneumonia^{2,3,9,10,12}

- •Review of the literature showed variable evidence for the ability to differentiate bacterial versus viral versus atypical pneumonia based on clinical, radiographic, or laboratory findings with a lack of clear predictive factors.
- •Patients with atypical pneumonia are typically older (> 4 years) and may more frequently demonstrate extrapulmonary findings such as rash or GI involvement.
- •The presence of wheezing may be more suggestive of a viral etiology.
- •Normal procalcitonin values may be less suggestive of a bacterial etiology. Elevated levels can be seen in bacterial, viral, and atypical causes. Routine labs are not recommended for mild to moderate pneumonia.

Appendix C: Signs that patient is not tolerating work of breathing

- •Inability to coordinate suck/swallow/breathing for feeds
- •Work of breathing prohibiting activities of daily living (eating, walking, talking)
- Persistent tachycardia despite adequate fluid resuscitation
- Change in mental status (inconsolable, lethargic)

Appendix D: Considerations for direct admit to Acute Care hospitalization

- For questions regarding management or hospitalization at the Milwaukee campus, please contact the Physician Consult Center (414-266-2470)
- For questions regarding management or hospitalization in Fox Valley, please page the FV Peds Hospitalist team ((920) 554-4502 need to confirm)
- Direct Admit to CW main campus acute care vs ED transfer consider the following:
 - Patient must have one or more of the <u>moderate/acute care criteria</u> and <u>none of the severe critical care</u> criteria to be appropriate for direct admission to acute care
 - Referring provider must be able to titrate O2 to assess if meeting Acute vs Critical Care Criteria
 - Patient must not require rapid fluid resuscitation

Appendix E: Definition of Treatment Failure 11, 14

- •Lack of clinical improvement <u>after 72h</u> of appropriate antimicrobial therapy with appropriate antibiotic/dosing/frequency
- •Concern if persistent fevers, hypoxia, not tolerating work of breathing, elevated inflammatory markers (labs not routinely recommended for uncomplicated CAP) after 72 hours of appropriate therapy
- •Cough and radiographic findings of CAP can persist and should not be used in isolation to indicate concern for treatment failure
- •See antimicrobial considerations for recommendations if concern for treatment failure
- •If questions regarding antimicrobial therapy, page/Voalte Antimicrobial Stewardship. For holidays/after-hours, page the on-call ID provider
- •If questions regarding need for airway clearance or alternative pulmonary diagnosis, page pulmonary

For questions concerning this work, contact cappathway@childrenswi.org

Appendix F: Azithromycin^{2,4,5,13}

- Antibiotics may not be needed for patients with mild atypical pneumonia because M. pneumoniae infections
 are frequently self-limited. Factors such as community prevalence and severity of symptoms can be
 considerations in antibiotic treatment decisions
- Azithromycin could be considered to cover for atypical pneumonia in hospitalized patients who are not improving after 72 hours of IV ampicillin and clinical concern for atypical pneumonia
- Mycoplasma testing can be considered for hospitalized patients if clinical concern. Testing is not recommended for outpatients
- If antibiotics indicated for bacterial pneumonia, it is more important to prescribe amoxicillin. 40% of strep pneumonia is resistant to azithromycin for outpatient/EDTC isolates
- There is lack of evidence supporting routine use of azithromycin. Symptomatic M. pneumoniae infection is often self-limited with insufficient evidence to support the efficacy of antibiotics including azithromycin.
- There is lack of evidence regarding the effect of treatment on transmissibility. Treatment may be appropriate for individuals who have vulnerable close contacts based on the provider's clinical judgment.
- M. pneumoniae is frequently detected in asymptomatic patients
- Azithromycin has a prolonged serum elimination half-life and prolonged exposure to organisms on the respiratory tract mucosa. This has been associated with the selection of resistant organisms on mucosal surfaces of treated patients and may represent a source of resistant organisms to others in the community, including resistance to other bacteria (e.g. gram negatives) and to other antibiotics. The risk of obesity increases with even a single exposure to macrolides due to significant impacts on the microbiome.

Appendix G: Bronchial hygiene protocol

- To order, enter Protocol Initiation HOSP RCS-Bronchial hygiene. It is part of the GEN PEDS Pneumonia Admission order set
- The bronchial hygiene protocol consists of assessment and treatments based on the bronchial hygiene protocol
 score with the indication of: impaired mucociliary clearance, to treat clinically significant atelectasis, to improve
 an ineffective cough, to remove excessive secretions, and monitor patients that receive bronchial hygiene
 treatments at home. The protocol allows the Respiratory Care Provider to up-regulate and down-regulate the
 following treatments based on the patients score: Various positive expiratory pressure therapies, highfrequency vest/IPV, manual chest physiotherapy, mechanical cough assist, and basic bronchial hygiene therapy
 (incentive spirometry, ambulation, cough and deep breathing)

Appendix H: Penicillin allergy

- See Connect for additional information regarding oral penicillin challenge
- See Connect for additional information regarding beta-lactam side chain reactivity
- **High-risk**: Anaphylaxis; swelling (face, lips, throat); difficulty breathing; wheezing; skin peeling; mouth blisters; drop in blood pressure; syncope; seizures; serum sickness; fever; within one hour of medication: abdominal pain, rash, itching, multiple episodes of vomiting
- **Low-risk:** Rash or itching >1 hour from med administration, single episode of vomiting within an hour of med administration, dizziness, nausea, cough
- Non-allergic symptoms (amoxicillin/ampicillin would be appropriate in these cases): Runny nose, diarrhea, headache, vomiting with med administration, family history of allergy. Can offer penicillin challenge in these situations if family requests though not required.

Appendix I: Cefdinir

Original: 11/2024

• See Antimicrobial Stewardship Connect page on why cefdinir is not recommended for the treatment of pneumonia.

For questions concerning this work, contact

College of Wisconsin, Inc See Medical Disclaimer

References:.

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Revised: 4/2025 Process owner: CAP Pathway Team

Version History and Summary of Changes

- Version 1.0 (12/2/2024) Go live
- Version 1.1 (12/27/2024) Updated max daily dosing for ampicillin and added clarification on management of treatment failure patients.
- Version 1.2 (2/5/2025) Added contact method for reaching Antimicrobial Stewardship team
- Version 1.3 (3/11/2025) Revised wording on high vs low risk allergy vs non-allergic symptoms
- Version 1.4 (4/9/2025) Revised wording on contact method for reaching Antimicrobial Stewardship team. Added guidance for management of concurrent sinusitis

Medical Disclaimer

This Clinical Practice Guideline (CPG) is designed to provide a framework for evaluation and treatment. It is not intended to establish a protocol for all patients with this condition, nor is it intended to replace a clinician's judgement. Adherence to this CPG is voluntary. Decisions to adopt recommendations from this CPG must be made by the clinician in light of available resources and the individual circumstances of the patient. Medicine is a dynamic science; as research and clinical experience enhance and inform the practice of medicine, changes in treatment protocols and drug therapies are required. The authors have checked with sources believed to be reliable in their effort to provide information that is complete and generally in accord with standards accepted at the time of publication. However, because of the possibility of human error and changes in medical science, neither the authors nor Children's Hospital and Health System, Inc., nor any other party involved in the preparation of this work warrant that the information contained in this work is in every respect accurate or complete, and they are not responsible for any errors in, omissions from, or results obtained from the use of this information.

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