The Rise and Fall of the Z-Pak: Updates in Antibiotic Guidelines for Common Urgent Care Conditions Worth Knowing

Joshua Russell, MD, MSc, ELS, FCUCM, FACEP

Senior Editor - EM:RAP, UCMax Podcast Editor-in-Chief, The Journal of Urgent Care Medicine (JUCM) Legacy-GoHealth Urgent Care Metro Portland, Oregon



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Disclosure

I have no financial interests or relationships to disclose.



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October 03, 2024

Intermountain Health Receives Major Funding as Part of National Initiative to Advance Care for Children by Improving Antibiotic **Prescribing Practices for Kids with Common Respiratory Tract** Infections





(oot't know about you, but fin tilled of being the laked at about antibiotic stewardship with all realize it's a problem. And through it's common practice to blame urgent care providers for the situation, we aren't uniquely cuipable for artibiotic oversee.

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Intermountain Health has been approved for a funding award by the Patient-Centered Outcomes Research Institute (PCORI) to help advance care for children by improving antibiotic prescribing practices for acute respiratory tract infection throughout its multi-state service

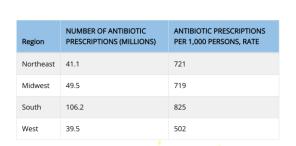
The \$2.5 million project will help Intermountain clinicians expand research-based practices for prescribing antibiotics to treat acute respiratory tract infection in children, avoid adverse side effects, and shrink the growing problem of drug-resistant infections.



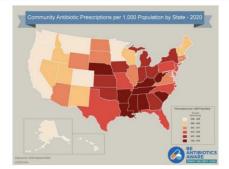
Learning Objectives

- Understand recent changes away from azithromycin in antibiotic guidelines for treatment of common bacterial infections encountered in Urgent Care
- 2. Develop familiarity with the few remaining indications of azithromycin
- 3. Identify the common and less common, but dangerous, adverse reactions associated with azithromycin

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1. Antibiotic prescriptions per 1000 persons by state (sextiles) for all ages — United States,



Regional Outpt Antibiotic Prescribing

Source: CDC, 2022



Can we accurately gauge patient expectations?

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INFECTIOUS DISEASE/ORIGINAL RESEARCH

Antibiotic Use for Emergency Department Patients With Upper Respiratory Infections: Prescribing Practices, Patient Expectations, and Patient Satisfaction

Samuel Ong, MD Janet Nakase, MPH Gregory J. Moran, MD David J. Karras, MD Matthew J. Kuehnert, MD David A. Talan, MD EMERGEncy ID NET Study Group From Olive View-University of California Los Angeles Medical Center, Sylmar, CA (Ong, Nakase, Moran, Talan); Temple University School of Medicine, Philadelphia, PA (Karras); and the Centers for Disease Control and Prevention (Kuenhert).

Study objective: Physicians often prescribe antibiotics to patients even when there is no clear indication for their use. Previous studies examining antibiotic use in acute bronchits and upper respiratory infections have been conducted in primary care settings. We evaluate the factors that physicians in the emergency department (ED) consider when prescribing antibiotics (eg, patient expectations) and the factors associated with patient satisfaction.

Methods: Ten academic EDs enrolled adults and children presenting with symptoms consistent with upper respiratory infection. Enrolled patients were interviewed before their physician encounter and were reinterviewed before discharge and 2 weeks later. Physicians were interviewed about factors that influenced their management decisions, including their perceptions of patients' expectations. Patients with a single diagnosis of uncomplicated acute bronchitis or upper respiratory infection were included for analysis.

Results: 0f 272 patients enrolled, 68% of bronchitis patients and 9% of upper respiratory infection patients received antibiotics. Physicians were more likely to prescribe antibiotics when they believed that patients expected them (odds ratio (long 15.3; 9% confidence interval [cl] 2.9 to 9.6), although they were able to correctly identify only 27% of the patients who expected antibiotics. Satisfaction with the E0 visit was reported by 87% of patients who exceeded antibiotics. Satisfaction with the E0 visit was reported by 92% of patients who believed they had a better understanding of their illness but only by 72% of those who thought they had no better understanding (OR 4.4; 95% Cl 2.0 to 8.4).

Conclusion: Physicians in our academic EDs prescribed antibiotics to 68% of acute bronchitis patients and to fewer than 10% of upper respiratory infection patients. Physicians were more likely to prescribe antibiotics to patients who they believed expected them, although they correctly identified only about 1 in 4 of those patients. Platient satisfaction was not related to receipt of antibiotics but was related to the belief they had a better understanding of their illness. [Ann Emerg Med. 2007;50:213-220.]



Cognitive Costs of Abx Stewardship

JAMA Intern Med. 2014 December; 174(12): 2029-2031. doi:10.1001/jamainternmed.2014.5225.

Time of Day and the Decision to Prescribe Antibiotics

Jeffrey A. Linder, MD, MPH, Jason N. Doctor, PhD, Mark W. Friedberg, MD, MPP, Harry Reyes Nieva, BA, Caroline Birks, MD, Daniella Meeker, PhD, and Craig R. Fox, PhD Division of General Medicine and Primary Care, Brigham and Women's Hospital, Boston,

Results

There were 21 867 ARI visits to 204 clinicians in 23 practices that met our inclusion criteria; 44% resulted in antibiotic prescriptions (Table). Antibiotic prescribing increased throughout the morning and afternoon clinic sessions for antibiotics sometimes indicated and antibiotics never indicated ARIs (Figure). Relative to the first hour of a session, the adjusted odds ratios of antibiotic prescribing in the second, third, and fourth hours were 1.01 (95% CI, 0.91–1.13), 1.14 (95% CI, 1.02–1.27), and 1.26 (95% CI, 1.13–1.41), respectively (P < .001 for linear trend).





Automating the Common & Difficult Conversation

SCRIPTING YOUR LEAST FAVORITE ANTIBIOTICS FOR

Courtesy of Rob Orman, MD

(www. roborman.com/free-resources)

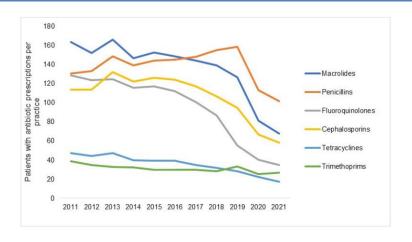
- Revise & Improve with Practice
- Save mental & emotional energy for MDM
- Start with top 2-3 most common painful conversations

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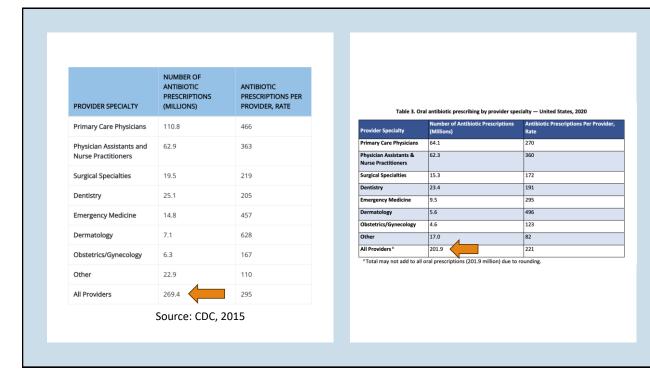


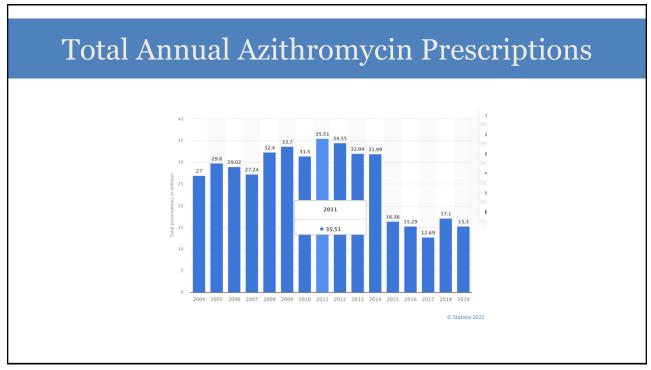
Did COVID Usher in a New Era of Antibiotic Stewardship?

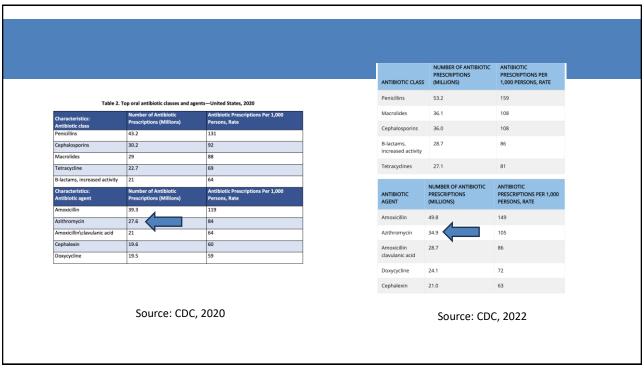
COVID-19 & Antibiotic Utilization

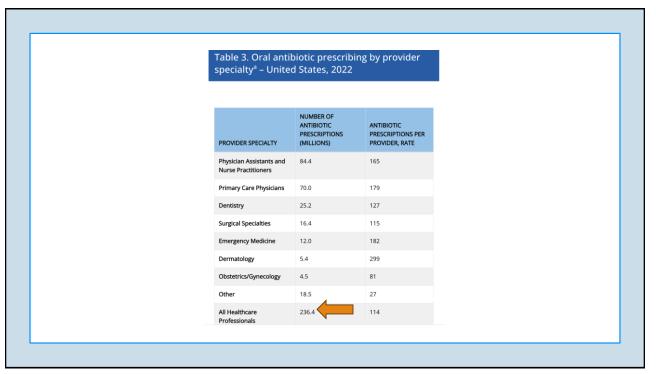


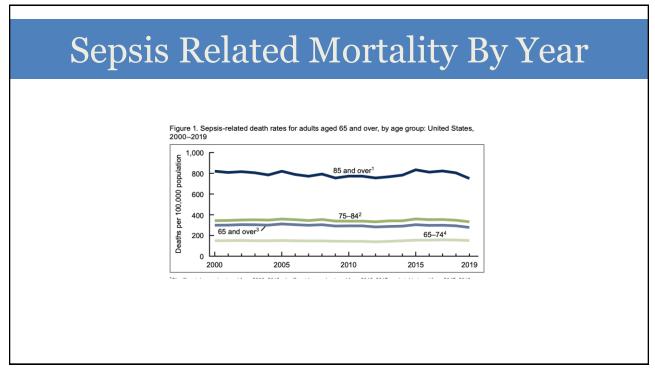
Tanislav C, Rosenbauer J, Kostev K. The COVID-19 Pandemic Enhanced the Decade-Long Trend of the Decreasing Utilization of Antibiotics. *Antibiotics*. 2023; 12(5):927.



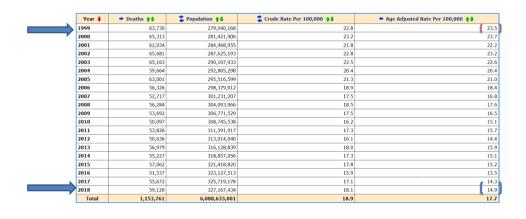








Pneumonia Related Mortality by Year





The Rise of the Z-Pak...

- Synthetic Macrolide developed by Pliva Pharmaceuticals, Yugoslavia in 1981
- Licensed to Pfizer in 1991 Z-Pak is born...
- Activity at 50S Ribosomal Subunit
- Clinically Favorable Profile
 - Delivered by phagocytes
 - Active at low pH
 - Long tissue half-life
- · Rapidly became among top 5 most commonly prescribed antibiotics...

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Azithromycin Resistance

- Population Level:
 - Rapid Emergence of Resistance of *S. pneumoniae* (& *S. pyogenes*) in 90's
 - Ribosomal Methylation (erm) or Efflux Pumping (mef)
 - Range of S. PNA Resistance from 10 -> 90% based on region

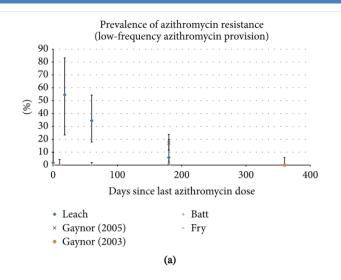


 Asymptomatic Pneumococcal Carriage Rates up to 90%



Pneumococcal Resistance

Derek K-H. Ho et al, "Antibiotic Resistance in *Streptococcus pneumoniae* after Azithromycin Distribution for Trachoma", *Journal of Tropical Medicine*, vol. 2015, Article ID 917370, 8 pages, 2015. https://doi.org/10.1155/2015/917370

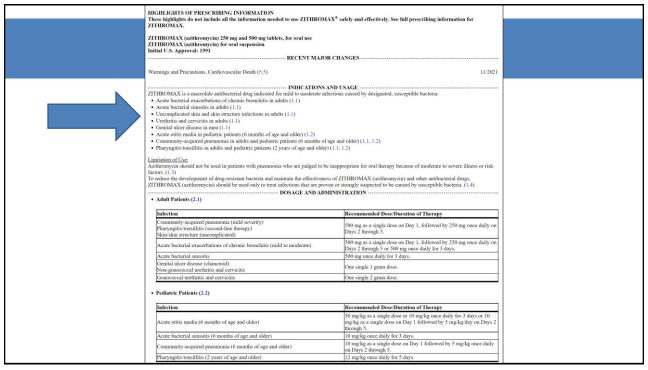


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"Gimme a Z-Pak to Knock It Down"

Common Z-Pak Requests:

- Sinusitis
- Cough/Bronchitis
- AOM in PCN allergy
- Strep in PCN allergy
- Chlamydia



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Acute Bacterial Sinusitis (ABRS)

Acute Sinusitis: Inflammation of nasal cavity & paranasal sinuses for <4 weeks

ABRS definition (per IDSA):

- 1. 10 days of illness w/o improvement
- 2. Severe symptoms (facial pain, purulent discharge) <u>AND</u> Fevers >39C x 3-4d
- 3. 'Double sickening' (new onset fever, facial pain, headache after URI)
- <2% of cases are bacterial & 80% of ABRS resolves w/I 2 weeks w/o ABX!!
- <u>Higher Risk for Complicated Course</u>: Advanced age, diabetes, immunosuppression

ABRS: What Do the Guidelines Say?



AAO-HNS (2015) | IDSA (2012) | ACP- CDC High Value Care Task Force (2016)

- 1. Treat only ABRS
- 2. Treat Immediately (IDSA) or Watch & Wait x 7 days (AAO-HNS)
 - Watchful waiting only if immunocompetent and good follow-up
- 3. "Azithromycin/macrolides NOT recommended for empiric therapy due to high rates of S. pneumoniae resistance"

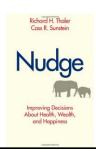
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ABRS: What Do the Guidelines Say?

- Amoxicillin/Clavulanate 875mg BID (NOT amoxicillin alone Resistance in H. flu and M. catarrhalis)
- Doxycycline 100mg BID (PCN allergy/alternate first line)
- Cefpodoxime 200mg BID

**Levofloxacin 500mg daily (only if can't tolerate alternatives; FQ risks)

Duration: 5-7 days



Community-Acquired Pneumonia (CAP)

Clinical diagnosis with constellation of findings: fever, dyspnea, cough, sputum production, abnormal lung sounds, abnormal CXR findings

- 'Clinicians should not perform testing or initiate antibiotic therapy in patients with bronchitis unless pneumonia is suspected' -ACP/CDC High Value Care Task Force, 2016
- Remember to educate: Up to 3-4 weeks of cough is COMMON with bronchitis
- Azithromycin is NOT recommended for outpatients (or inpatients) with COVID-19



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CAP: What Do the Guidelines Say?

2019 ATS/IDSA Joint Guideline Updates on Treatment of CAP:

- **Amoxicillin 1g TID -OR- Doxycycline 100mg BID x 5 days** (healthy, <65yo w/o recent antibiotic use)
- Azithromycin/Macrolide ONLY Recommended if Local S. Pneumoniae resistance <25%
- Dual Therapy for >65 and/or co-morbidities and/or recent antibiotic use:
 - Amox/clav -OR- 3rd gen cephalosporin PLUS Doxycycline -OR- Macrolide
 - ** Monotherapy with respiratory fluoroquinones (e.g. levofloxacin) least preferrable option

Penicillins in Urgent Care

Strep Pharyngitis

- Preferred first line: Penicillin VK (or Amoxicillin)
- Macrolide resistant S. pyogenes

Acute Otitis Media

- Preferred first line: Amoxicillin (+/- Clavulanate)
- Macrolide resistant S. pneumoniae
- Use WAIT-AND-SEE more

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"I'm Allergic to Penicillin"

Penicillin is the most commonly reported drug allergy.¹





of patients in the US report penicillin allergy.¹ 9 out of 10 reporting penicillin allergy are not truly allergic.4





80% of patients with IgE-mediated penicillin allergy lose the sensitivity after 10 years.4

Au LYC, Siu AM, Yamamoto LG. Cost and Risk Analysis of Lifelong Penicillin Allergy. *Clin Pediatr (Phila)*. 2019;58(11-12):1309-1314. doi:10.1177/0009922819853014

Consequences of 'Penicillin Allergy'

- Higher Lifelong Healthcare Spending
- Higher Rates of Broad Spectrum & Quinolone Abx Exposure
- Higher Rates of C. difficile



Au LYC, Siu AM, Yamamoto LG. Cost and Risk Analysis of Lifelong Penicillin Allergy. *Clin Pediatr (Phila)*. 2019;58(11-12):1309-1314. doi:10.1177/0009922819853014

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Guidelines: De-Label When Able

Penicillin Risk Assessment

- Adverse Reaction/Intolerance (e.g. GI upset) → PCN and Amoxicillin Safe
- 2. Mild/Delayed Hypersensitivity (e.g. maculopapular rash) → PO Test Dose of PCN or Cephalosporin
 - Cephalosporin/Penicillin Cross Reactivity (mostly) Myth
 - 97+% w/ true PCN allergy tolerate cephalosporins
 - · Consider Allergist Referral for Formal Testing
- 1. Immediate/**IgE** Mediated (e.g. hives, anaphylaxis) → 3rd or later generation cephalosporin preferrable, but all likely safe (0.001% risk of anaphylaxis)
- 2. Severe Non-Allergic Reaction (e.g. TEN, SJS, DRESS) → Avoid ALL B-lactams

De-Labeling Guidelines

Chua, KY et al. "The Penicillin Allergy Delabeling Program: A Multicenter Whole-of-Hospital Health Services Intervention and Comparative Effectiveness Study." Clinical infectious diseases: an official publication of the Infectious Diseases Society of America vol. 73,3 (2021): 487-496.

Dermatological				Respiratory or Systemic		Unknown								
Skin manifestation			commendation & ultant allergy type			commendation & ultant allergy type	Clinical manifestation	Clinical manifestation Recommendation & Resultant allergy type						
Childhood exanthem (unspecified) Mild rash with no severe features			Unlikely to be significant (non-severe)	Laryngeal involvement ("throat tightness" or "hoarse voice")	Immediate hypersentitisty (severe)		Unknown reaction ≤ 10 years ago		Unlengum (non-severe)					
Immediate diffuse rash ("itchy immediate rash") <2 hours post dose		0	Immediate hypersensitivity (non-severe)			Unknown reaction > 10 years ago or family history of penicillin allergy only	0	Unlikely to be significant (non-severe)						
Diffuse rash or localized ago or unknown			Delayed hypersensitivity (non-severe)	Respiratory compromise ("shortness of breath")		Immediate hypersensitivity (severe)	Renal							
with no other symptoms (non-immediate or unknown timing)	≤ 10 years ago		Delayed hypersensitivity (non-severe)	Fever ("high temperature") Not explained by infection		Delayed hypersensitivity (severe)	Severe renal injury, failure or AIN >50% reduction in eGFR from baseline or absolute serum creatinine increase of ≥26.5µmol/L, or transplantation, or dialysis		Potential immune mediated (severe)					
Angioedema ("lip, facial or tongue swelling")		0	Immediate hypersensitivity (severe)	Anaphylaxis or unexplained collapse		Immediate hypersensitivity (severe)	Mild renal impairment (Does not meet criteria in box above)	_	Unlikely immune mediated (non-severe)					
Generalized swelling (outside of angioedema)			Immediate hypersensitivity (severe)	Haematological		Liver								
Urticaria ("wheals and hives")						termediate hypersensitivity	hypersenstivity		Low platelets < 150 x10°/L or unknown	_	Potential immune mediated (severe)	Severe liver injury, failure or DILI (£5x upper limit of normal (ULN) for ALT or AST, or ≥3x ULN for ALT with ≥2x ULN for bilirubin, or ≥2x ULN for ALP, or transplant)	_	Potential immune mediated (severe)
			(non-severe)	Low neutrophils < 1x10 ⁹ /L or unknown		Potential immune mediated (severe)	Mild hepatic enzyme derangement (Does not meet criteria in box above)		Unlikely immune mediated (non-severe)					
Mucosal ulceration ("mouth, eye or genital ulcers")			Delayed hypersensitivity (severe)	Low haemoglobin < 100 g/L or unknown		Potential immune mediated (severe)	Gastrointestinal, Neurological or Infusion-related		Infusion-related					
Pustular, blistering or desquamating rash ("skin shedding")			Delayed Impersensitivity	Eosinophilia (>0.7 x 10 ⁹ /L or unknown)		Delayed hypersensitivity (severe)	Gastrointestinal symptoms ("nausea, vomiting, diarrhoea")		Unlikely immune mediated (non-severe)					
			(severe)				Mild neurological manifestation ("headache, depression, mood disorder")	_	Unlikely immune mediated (non-severe)					
Appropriate for supervised direct oral rechallenge (or direct de-labelling)					□ Low risk	Severe neurological manifestation		Unknown or unclear						
Appropriate for supervised direct oral rechallenge					□ Low risk	("seizures or psychosis")		mechanism						
May be appropriate for referral for specialized skin testing					□ Moderate risk	Anaphylactoid/infusion reaction		Unknown or unclear						
May be appropriate for referral for specialized skin testing					□ High risk	(e.g. red man syndrome)		mechanism						

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Penicillin Allergy Risk: 'PEN-FAST'

PEN	Penicillin allergy reported by patient	[] If yes, proceed with assessment
F	Five years or less since reaction ^a	2 points
A	Anaphylaxis or angioedema	2 points
S	Severe cutaneous adverse reaction ^b	
т	Treatment required for reaction ^a	1 point
		Total points
	Interpretation	
Points 0 Ver	y low risk of positive penicillin allergy test <1	% (<1 in 100 patients reporting penicillin allergy)
1777	risk of positive penicillin allergy test 5% (1 in	20 patients)
1-2 Lov		
ii	derate risk of positive penicillin allergy test 20	% (1 in 5 patients)

Trubiano JA, Vogrin S, Chua KYL, et al. Development and Validation of a Penicillin Allergy Clinical Decision Rule. *JAMA Intern Med.* 2020;180(5):745-752. doi:10.1001/jamainternmed.2020.0403

PEN-FAST in Practice

- 73% of patients had PEN-FAST 0-2
- No patient with PEN-FAST <3 had +skin test
- 13% had safe Direct Oral Challenge

Performance Measure	Cutoff PEN-	Cutoff PEN-FAST score				
	0 vs 1-5	0-1 vs 2-5	0-2 vs 3-5			
Sensitivity, % (95% CI)	100 (39.8-	100 (39.8-	100 (39.8-			
	100)	100)	100)			
Specificity, % (95% CI)	18.1 (11.6-	69.8 (60.6-	75.9 (67.0-			
	26.3)	78.0)	83.3)			

Su C, Belmont A, Liao J, Kuster JK, Trubiano JA, Kwah JH. Evaluating the PEN-FAST Clinical Decision-making Tool to Enhance Penicillin Allergy Delabeling. *JAMA Intern Med*. 2023;183(8):883-885. doi:10.1001/jamainternmed.2023.1572

Penicillin skin testing, No. (%)	104 (86.7)
Oral challenge, No. (%)	118 (98.3)
Oral challenge following skin test	102 (85.0)
Direct oral challenge	16 (13.3)
PEN-FAST score	
0	21 (17.5)
1	60 (50.0)
2	7 (5.8)
3	30 (25.0)
4	0
5	2 (1.7)

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PEN-FAST Limitations

- Not Prospectively Validated in Adults (Bias?)
- **Higher Risk Features:** Rapid Onset of Reaction, Short Time Since Last Reaction, Requiring Hospitalization
- Not Valid in Children (i.e. < 18 years)

Copaescu AM, Vogrin S, Shand G, Ben-Shoshan M, Trubiano JA. Validation of the PEN-FAST Score in a Pediatric Population. *JAMA Netw Open*. 2022;5(9):e2233703

PEN-FAST - My Practice

- Don't use in Pts <18 yrs
- Use Shared Decision Making
- Direct Oral Challenge in Clinic if PF = 0-1
- If 2 or greater, refer to allergist
- Cephalosporins Generally Safe

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Wait-and-See for AOM

- RCT of 283 children 6m 12y seen in ED
- Abx filled: Immediate 62 % vs WASP 13%
- No difference in subsequent fever, otalgia, or revisit

Wait-and-See Prescription for the Treatment of Acute Otitis Media

A Randomized Controlled Trial

Khoon-Yen Tay, MD Donald H. Arnold, MD, MPH James D. Dziura, PhD Mark D. Baker, MD

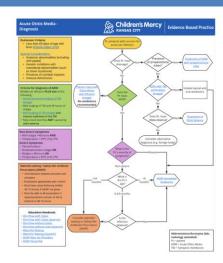
CUTE OTITIS MEDIA (ACM) is the most common reason for which an antibotic is many the most common reason for which an antibotic is a missing of the most common reason for which an antibotic is a missing of the most common reason is a missing or most for an extract 15 million antibotic precipitions written per year in the United States. *Untracted ACM has a high an attention of ACM accounts for an extract of spontaneous resolution, with similar of spontaneous resolution, with similar arts of complications whether arts of spontaneous resolution, with similar arts of spontaneous resolution, with similar arts of spontaneous resolution, with similar of spontaneous resolution, with similar to spontaneous resolution, with similar of spontaneous resolution in the subject of the second CUTE OTITIS MEDIA (AOM) IS

Context Acute otilis media (AOAN) is the most common diagnosis for which antib otics are prescribed for children. Previous risals that have evaluated a "walt-and-se prescription" (WASP) for antibiotics, with which parents are asked not to fill the pre-scription "WASP) for antibiotic on the previous results of the previous pre

Objectives To determine whether treatment of AOM using a WA reduces use of antibiotics compared with a "standard prescription" (SP) and to eval ate the effects of this intervention on clinical symptoms and adverse outcomes relat to antibiotic use.

Wait-and-See Criteria

- 1. Non-severe sxs*
- 2. >6 months age
- 3. Unilateral
- 4. Can f/u w/i 2-3 days



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Chlamydia trachomatis - most common bacterial sexually transmitted genital infections

2nd Most Common Reportable Disease & 20% Inc since 2015

Chlamydia

Intracellular Reproduction

Presentation ranges from Asymptomatic -> Dysuria/Discharge -> PID

Chlamydia: What Do The Guidelines Say?

CDC STI Treatment Guidelines, 2021

- Doxycycline 100mg BID x 7 days (1st line)
- 20% Failure Rate w/ Azithromycin 1g PO
- Doxy more effective for rectal and pharyngeal chlamydia infection
- Azithromycin 1g PO x 1 still preferred in pregnancy or if expect non-adherence with doxy
 - Consider "test of cure"

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Toxicity and Adverse Reactions

- <u>Black box</u>: "Rare QTc prolongation and ventricular arrhythmias, including torsades de pointes"
- Gastrointestinal Immediate (vomiting) & Delayed (diarrhea)
- Drug-Drug Interactions (CYP3A4)
- Liver Injury (rare, usually reversible, but fatal in cases if not stopped)



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When might Azithromycin be indicated?

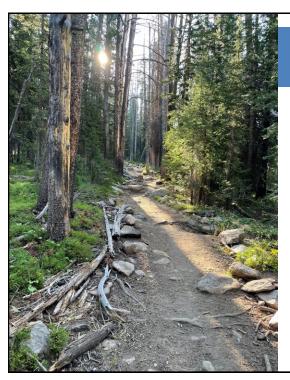
- 1. Traveler's diarrhea
- 2. COPD Exacerbation
- 3. Atypical pneumonia
- 4. Chlamydia in Pregnancy or ?Adherence



Drug	Age	Dose	Duration	Other Considerations
Rifampin	<1 month	5 mg/kg, orally, every 12 hours	2 days	Consider consultation with a pediatric infectious disease expert for infants <1 month.
	≥1 month	10 mg/kg (maximum 600 mg), orally, every 12 hours	2 days	Can interfere with efficacy of oral contraceptives and some seizure prevention and anticoagulant medications; may stain soft contact lenses. Not recommended for pregnant people.
Ceftriaxone	<15 years	125 mg, intramuscularly	Single dose	To decrease pain at injection site, dilute with 1% lidocaine.
	≥15 years	250 mg, intramuscularly	Single dose	
Azithromycin	All Ages	10 mg/kg (maximum 500 mg)	Single dose	Alternative agent. In one study, equivalent to rifampin for eradication of N . meningitidis from nasopharynx.

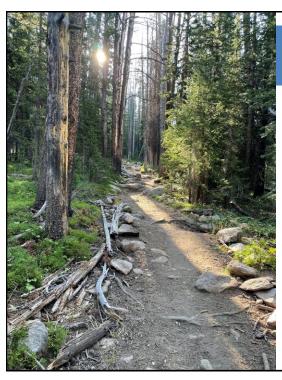
Meningococcal Prophylaxis?

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SUMMARY

- ☐ Patient satisfaction poorly correlated w/ Abx Rx
- ☐ Azithromycin stewardship improving...but still overprescribed in UC
- \square S. PNA & S. Pyogenes resistance is common



SUMMARY

- ☐ Azithro never recommended for ABRS
- ☐ Azithro monotx is out for CAP
- ☐ PCN "allergy" can usually be de-labeled
- ☐ Wait-and-See for AOM
- ☐ Doxy is the way to go for Chlamydia
- ☐ Introduce the 'Fall' of Azithromycin to explain selflimited illness and risks