



Noam was running a high fever and complained of a **painful throat** that made it difficult to eat



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Group A β -hemolytic streptococcus and Tonsillopharyngitis

GABHS or *Streptococcus pyogenes* is the most common bacterial cause of acute pharyngitis in children¹⁻³

Among children of all ages who present with sore throat, the prevalence of **GABHS** has been estimated to be **37%**^{2,4*t}

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GABHS tonsillopharyngitis: Treatment Goals



- In tonsillopharyngitis, the primary outcome and antibiotic treatment goal of interest is eradication of **GABHS**^{3,5}
- **Eradication is necessary**⁵
 - To prevent non-suppurative and suppurative sequelae
 - To eliminate contagion
 - To produce a more rapid symptomatic resolution of the illness

GABHS: group A β -hemolytic streptococci

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Results of the Survey of Antibiotic Resistance



Susceptibility of *S. pyogenes* to Cefuroxime axetil*

	Country	No. of isolates	% Susceptibility ^a	Isolates collection period
	India ⁶	78	100	2012-14
	Turkey ⁷	222	100	2011-13
	Pakistan ⁸	95	100	2007-09

a: CLSI guidelines assume susceptibility to cephalosporins based on penicillin susceptibility.

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Cefuroxime axetil vs. Penicillin V in Group-A β -hemolytic Streptococcal Tonsillopharyngitis

Meta-analysis of randomized, controlled trials showed higher bacteriological and clinical cure rates with cefuroxime as compared to penicillin in treatment of group A β -hemolytic streptococcal (GABHS) tonsillopharyngitis in children.⁵

Analysis of bacteriological cure rate						
	Penicillin n/N	Favours Penicillin	OR 1	Favours Cefuroxime	Cefuroxime n/N	OR (95% CI Random)
Data from 4 clinical trials	61/109				99/114	5.19(2.68, 10.07)
	106/126				244/259	3.07(1.51, 6.23)
	29/33				51/60	0.78(0.22, 2.26)
	30/38				75/77	10.00(2.01, 49.83)
Total	226/306				469/510	3.31(1.47, 7.45)

GABHS: group A β -hemolytic streptococci

Adapted from: Casey JR, Pichichero ME. Meta-analysis of cephalosporin versus penicillin treatment of group A streptococcal tonsillopharyngitis in children. *Pediatrics*. 2004;113(4):866-82.

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	Penicillin n/N	Favours Penicillin	OR 1	Favours Cefuroxime	Cefuroxime n/N	OR (95% CI Random)
Data from 4 clinical trials	73/109				98/114	3.02(1.56, 5.86)
	102/126				238/259	2.67(1.42, 5.01)
	29/33				60/60	18.46(0.96, 453.33)
	38/38				77/77	Not Estimable
Total	242/306				473/510	2.96(1.88, 4.64)

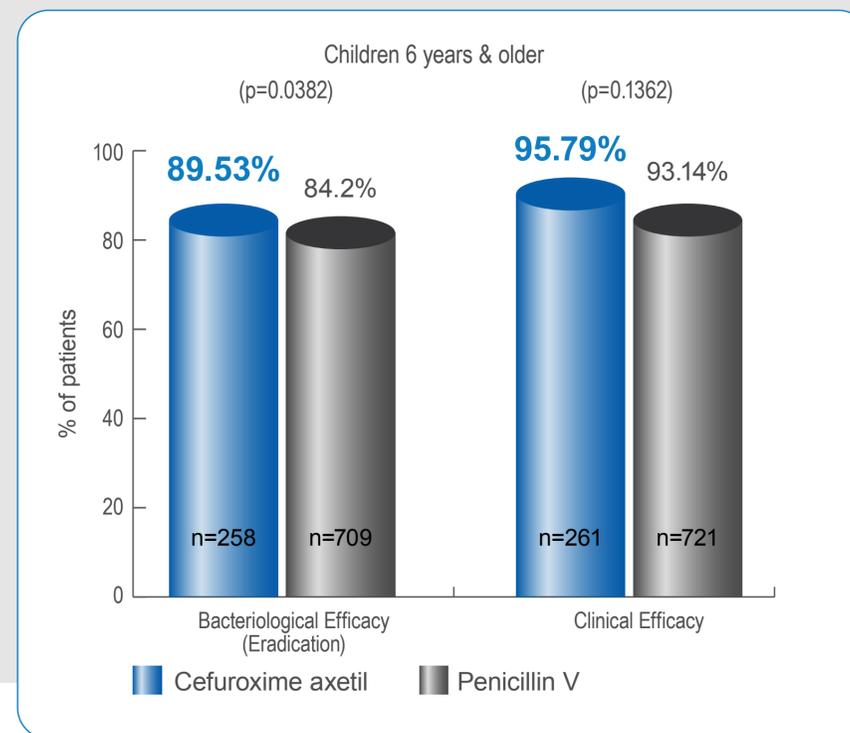
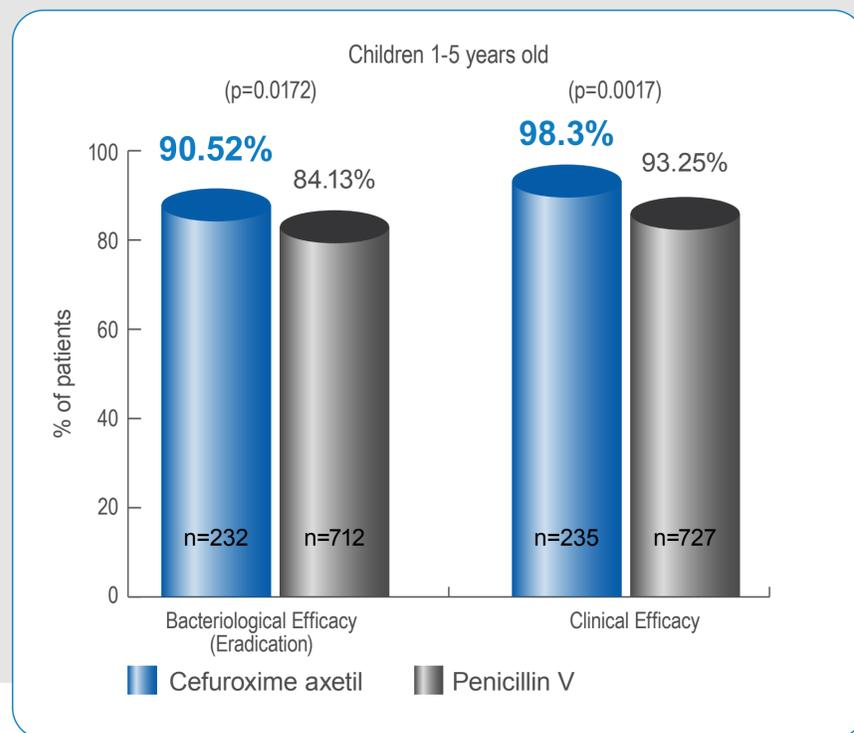
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Bacteriological and clinical efficacy rates

Cefuroxime axetil b.i.d. for 5 days was superior to oral Pen V t.i.d. for 10 days in eradicating GABHS⁹



Cefuroxime axetil:
20 mg/kg/day (max 500 mg) in
2 doses x 5 days.

Penicillin V: 50,000IU/kg
(30 mg/kg) daily in
3 doses x 10 days.

Shorter therapeutic course with
cefuroxime axetil as compared
to penicillin for GABHS
tonsillopharyngitis may improve
compliance in children.⁹⁻¹¹

Efficacy Rates 2-4 days after completion of treatment

GABHS: group A β-hemolytic streptococci

Adapted from: Scholz H. Streptococcal-A tonsillopharyngitis: a 5-day course of cefuroxime axetil versus a 10-day course of penicillin V. results depending on the children's age. *Chemotherapy*. 2004;50(1):51-4.

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Summary



- GABHS or *Streptococcus pyogenes** has high susceptibility to Cefuroxime axetil⁶⁻⁸
- Cefuroxime axetil has higher bacteriological and clinical cure rates as compared to penicillin in treating GABHS tonsillopharyngitis in children⁵
- Shorter therapeutic course with cefuroxime axetil as compared to penicillin for GABHS tonsillopharyngitis may improve compliance in children⁹⁻¹¹

*Susceptibility patterns may vary with time and geography.
Kindly refer to local susceptibility patterns.

GABHS: group A β -hemolytic streptococci

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Zinnat main safety information^{12,13}



Contraindicated in patients with known hypersensitivity to cephalosporin antibiotics, and special care must be taken in patients with previous allergic reaction to penicillins or other β -lactams.



Pseudomembranous colitis has been reported with the use of antibiotics and may range in severity from mild to life-threatening.



Prolonged use may result in overgrowth of *Candida* and other non-susceptible organisms (e.g. enterococci and *C. difficile*).

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Zinnat main safety information^{12,13}



The most commonly reported adverse drug reactions ($\geq 1/100$ to $< 1/10$) are headache, dizziness, eosinophilia, overgrowth of *Candida*, gastrointestinal disturbances including diarrhoea, nausea and abdominal pain; and transient increases of hepatic enzyme levels.



In patients with renal impairment the dosage will need to be adjusted.



Consideration should be given to local susceptibility data (where available) and official guidance on the appropriate use of antibacterial agents.

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Dosage and administration

The usual course of therapy is seven days (range five to ten days)^{12,13}



Tablet¹²

Cefuroxime axetil tablets should be taken after food for optimum absorption.



Adults and children (≥40 kg)

MOST INFECTIONS: 250 mg twice daily



Children (<40 kg)

MOST INFECTIONS: 15 mg/kg twice daily, to a maximum of 250 mg daily.

Acute tonsillitis and pharyngitis, acute bacterial sinusitis the recommended dose is 10 mg/kg twice daily to a maximum of 125 mg twice daily



Suspension¹³

For optimal absorption cefuroxime axetil suspension should be taken with food.

Adults and children (≥40 kg)

MOST INFECTIONS: 250 mg twice daily

Children (<40 kg)

The usual dose of Zinnat Suspension is 10 mg/kg (a maximum of 125 mg) to 15 mg/kg (a maximum of 250 mg) twice daily depending on the severity and type of infection and the weight and age of the child.

Acute tonsillitis and pharyngitis, acute bacterial sinusitis the recommended dose is 10 mg/kg twice daily to a maximum of 125 mg twice daily

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