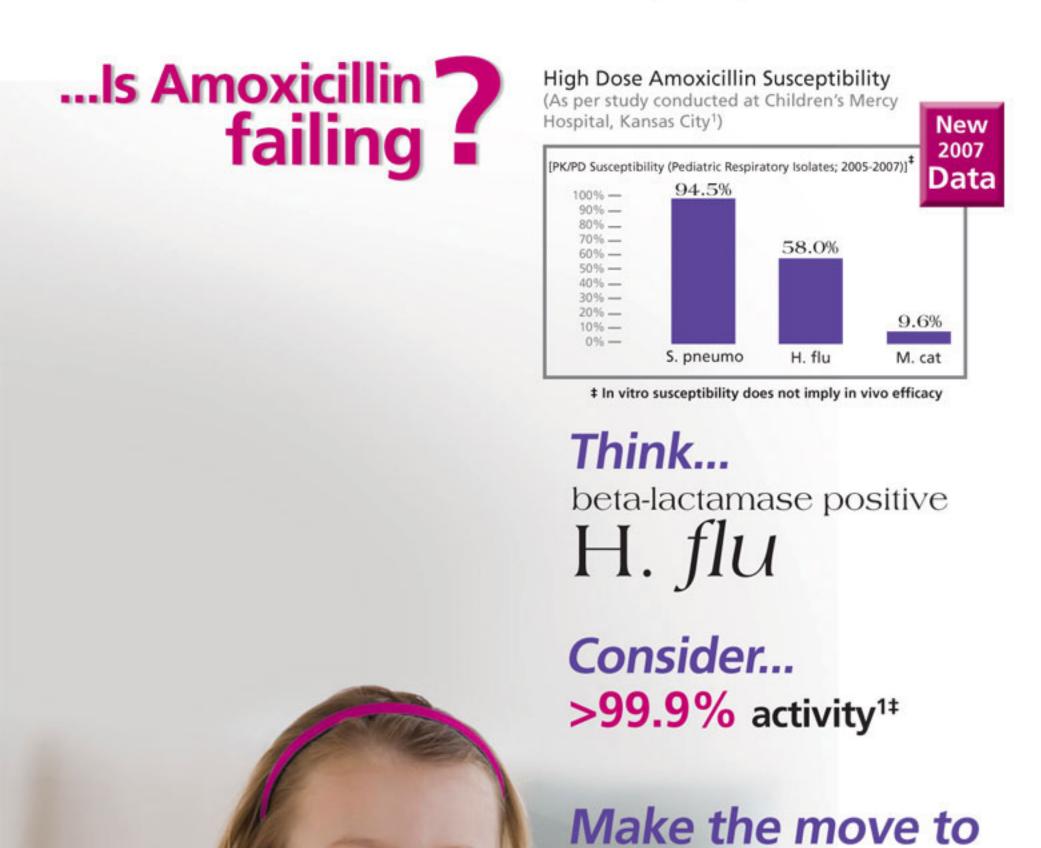
In acute otitis media (AOM)*...



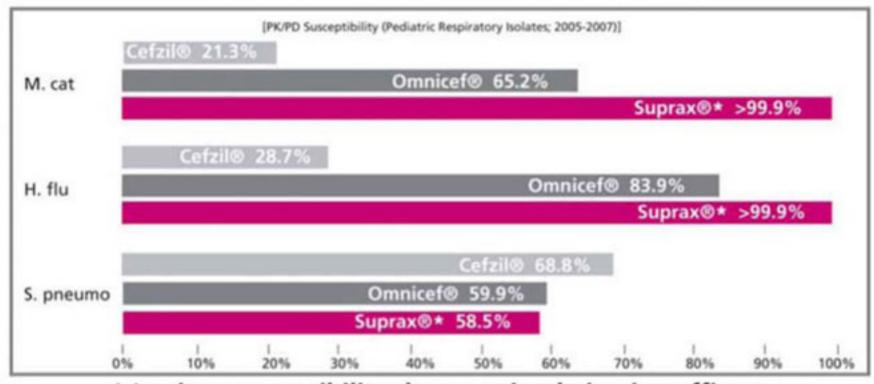
Once-a-day

200 mg/5 mL

Make the move to

More active against H. flu and M. cat, in vitro11

(As per study conducted at Children's Mercy Hospital, Kansas City)



‡ In vitro susceptibility does not imply in vivo efficacy

Offers greater activity and less resistance to H. flu than any other oral suspension^{2,3}

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*SUPRAX o is indicated for the treatment of otitis media caused by H. influenzae (B-lactamase +/- strains), M. (Branhamella) catarrhalis (most of which are B-lactamase positive), and S. pyogenes.

References: 1. Data on file; Lupin Pharmaceuticals, Inc.; 2007. 2. Sinus and Allergy Health Partnership. Ortolaryngol Head Neck Surg. 2004; 130(suppl): 1-45. 3. Adapted from Leibovitz E, Jacobs MR, Dagan R. Haemophilus influenzae: a significant patheogen in acute otitis media. Pediatr Infect Dis J. 2004; 23:1142-1155.

Please see attached full prescribing information.

SUPRAX® Tablets are not recommended for AOM.



Lupin Pharmaceuticals, Inc., Harborplace Tower, 111 South Calvert St., Suite 2150 Baltimore, MD 21202 LUPIN www.lupinpharmceuticals.com A Commitment to Caring for Kids

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Ordering information

The following item(s) are available for ordering from your Cardinal Health Customer Service team.

Description	Strength	Size	Form	NDC #	CIN
Suprax®	200 mg/5 mL	50 mL	Powder-Oral Suspension	27437-0206-03	3970225
Suprax®	200 mg/5 mL	75 mL	Powder-Oral Suspension	27437-0206-02	3970191
Suprax®	100 mg/5 mL	50 mL	Powder-Oral Suspension	68180-0202-03	3565728
Suprax®	100 mg/5 mL	100 mL	Powder-Oral Suspension	68180-0202-01	3565744
Suprax®	400 mg	50 ct	Tablet	27437-0201-08	4058897

Please visit www.lupinpharmaceuticals.com/supraxinfo.htm for full prescribing information.

Available in select Cardinal Health Distribution Centers. Please contact your Cardinal Health Customer Service team for more information.

- 1. Suprax® [package insert]. Baltimore, MD: Lupin Pharmaceuticals, Inc. 2004.
- Data on file, Lupin Pharmaceuticals, Inc.; 2004.

Important safety considerations: The most common adverse event related or probably related to SUPRAX® was GI upset. Several patients developed severe diarrhea and/or documented pseudomembranous colitis, and a few required hospitalization. SUPRAX® is contraindicated in patients with known allergy to the cephalosporin group of antibiotics.

WARNINGS: BEFORE THERAPY WITH SUPRAX® IS INSTITUTED, CAREFUL INQUIRY SHOULD BE MADE TO DETERMINE WHETHER THE PATIENT HAS HAD PREVIOUS HYPERSENSITIVITY REACTIONS TO CEPHALOSPORINS, PENICILLINS, OR OTHER DRUGS. IF THIS PRODUCT IS TO BE GIVEN TO PENICILLIN-SENSITIVE PATIENTS, CAUTION SHOULD BE EXERCISED BECAUSE CROSS HYPERSENSITIVITY AMONG B-LACTAM ANTIBIOTICS HAS BEEN CLEARLY DOCUMENTED AND MAY OCCUR IN UP TO 10% OF PATIENTS WITH A HISTORY OF PENICILLIN ALLERGY. IF AN ALLERGIC REATION TO SUPRAX® OCCURS, DISCONTINUE THE DRUG. SERIOUS ACUTE HYPERSENSITIVITY REACTIONS MAY REQUIRE TREATMENT WITH EPINEPHRINE AND OTHE EMERGENCY MEASURES, INCLUDING OXYGEN, INTRAVENOUS FLUIDS, INTRAVENOUS ANTIHISTAMINES, CORTICOSTEROIDS, PRESSIR AMINES AND AIRWAY MANAGEMENT, AS CLINICALLY INDICATED.