AMOXICILLIN 500 MG- amoxicillin capsule
Health Department, Oklahoma State

Amoxicillin 500 mg Pack

Precaustions from Manufacturer Package Insert

5.1 Anaphylactic Reactions

Serious and occasionally fatal hypersensitivity (anaphylactic) reactions have been reported in patients on penicillin therapy including amoxicillin. Although anaphylaxis is more frequent following parenteral therapy, it has occurred in patients on oral penicillins. These reactions are more likely to occur in individuals with a history of penicillin hypersensitivity and/or a history of sensitivity to multiple allergens. There have been reports of individuals with a history of penicillin hypersensitivity who have experienced severe reactions when treated with cephalosporins. Before initiating therapy with amoxicillin, careful inquiry should be made regarding previous hypersensitivity reactions to penicillins, cephalosporins, or other allergens. If an allergic reaction occurs, amoxicillin should be discontinued and appropriate therapy instituted.

5.2 Clostridium difficile Associated Diarrhea

Clostridium difficile associated diarrhea (CDAD) has been reported with use of nearly all antibacterial agents, including amoxicillin, and may range in severity from mild diarrhea to fatal colitis. Treatment with antibacterial agents alters the normal flora of the colon leading to overgrowth of C. difficile.

C. difficile produces toxins A and B which contribute to the development of CDAD. Hypertoxin-producing strains of C. difficile cause increased morbidity and mortality, as these infections can be refractory to antimicrobial therapy and may require colectomy. CDAD must be considered in all patients who present with diarrhea following antibacterial use. Careful medical history is necessary since CDAD has been reported to occur over 2 months after the administration of antibacterial agents.

If CDAD is suspected or confirmed, ongoing antibiotic use not directed against C. difficile may need to be discontinued. Appropriate fluid and electrolyte management, protein supplementation, antibiotic treatment of C. difficile, and surgical evaluation should be instituted as clinically indicated.

5.3 Development of Drug-Resistant Bacteria

Prescribing amoxicillin in the absence of a proven or strongly suspected bacterial infection is unlikely to provide benefit to the patient and increases the risk of the development of drug-resistant bacteria.

5.4 Use in Patients With Mononucleosis

A high percentage of patients with mononucleosis who receive amoxicillin develop an erythematous skin rash. Thus amoxicillin should not be administered to patients with mononucleosis.

Repackaging Label


24
17

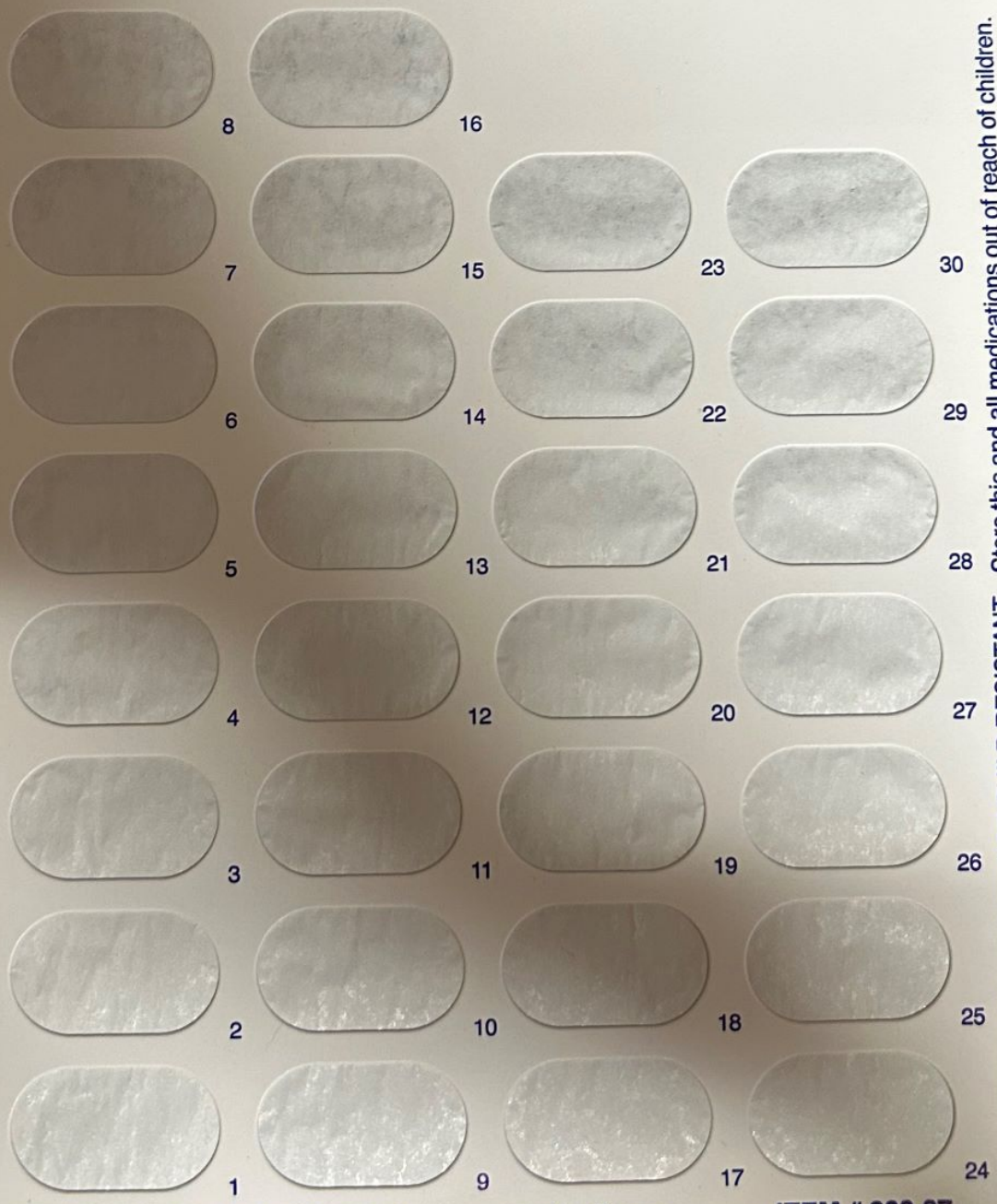
FILLED BY_____ CK'D BY_____
RECEIVED BY_____
START DATE_____
REORDERED BY_____
ORDER DATE_____

SURE Med® by Omnicell®

Packaged by:
OKLAHOMA STATE HEALTH DEPARTMENT OF HEALTH
7725 West Reno Avenue, Oklahoma City, OK 73127
405-426-8000

Amoxicillin 500 mg Capsule
Lot: FT5025069A
Expires: 12/07/26
MFG NDC: 57237-0031-01
Labeler Code: 83112-500-30

Store at 68 – 77°F (Excursions permitted 59-88°F)
Protect from light and moisture.
FEDERAL LAW PROHIBITS DISPENSING WITHOUT PRESCRIPTION



CAUTION: This package NOT CHILD RESISTANT. Store this and all medications out of reach of children.

MTS Medication Technologies®

omnicell.com

ITEM # 300-07

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START DATE_____
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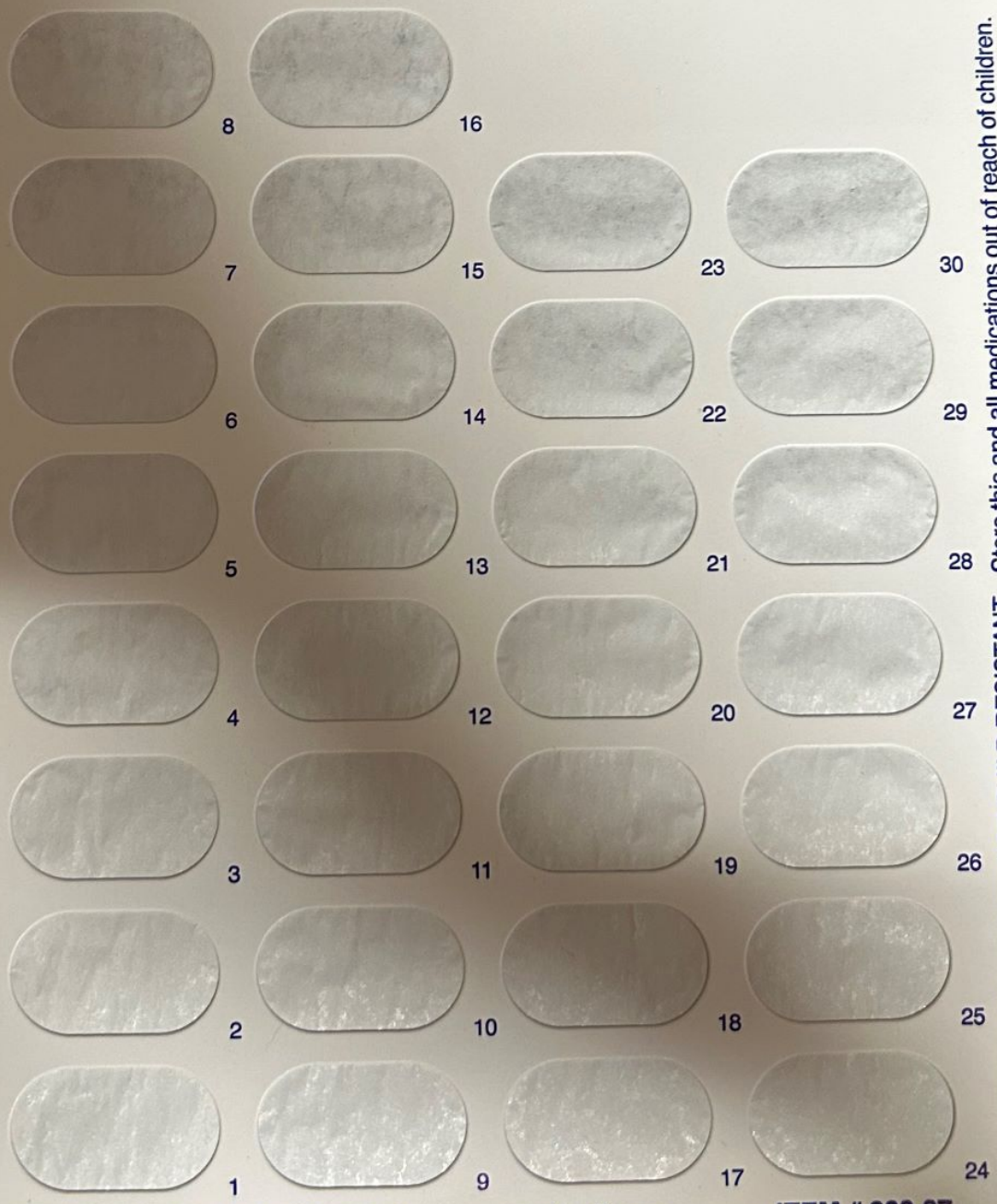

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ITEM # 300-07


PED
Amoxicillin 500mg
30 Count

AMOXICILLIN 500 MG

amoxicillin capsule

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:83112-500(NDC:57237-031)
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
AMOXICILLIN (UNII: 804826J2HU) (AMOXICILLIN ANHYDROUS - UNII:9EM05410Q9)	AMOXICILLIN ANHYDROUS	500 mg

Product Characteristics			
Color	blue, pink	**Score**	no score
Shape	CAPSULE	**Size**	23mm
Flavor		**Imprint Code**	A45
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83112-500-30	30 in 1 BLISTER PACK; Type 0: Not a Combination Product	11/09/2005	
2	NDC:83112-500-21	21 in 1 BLISTER PACK; Type 0: Not a Combination Product	11/09/2005	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA065271	11/09/2005	

Labeler - Health Department, Oklahoma State (143673015)

Revised: 12/2025 Health Department, Oklahoma State