ZYLAST XP ANTISEPTIC FOAMING- benzethonium chloride soap
Bocchi Laboratories Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Zylast Antiseptic Foaming Soap
Drug Facts

Active Ingredients

Benzethonium Chloride - 0.20%

Purpose

Antiseptic

Uses

- Handwash to decrease bacteria on the skin that potentially can cause disease.
- Recommended for repeated use.

Warnings

- For external use only.
- Avoid contact with eyes. In case of eye contact, rinse thoroughly with water.
- Discontinue use if irritation and redness develops. Consult a doctor if condition persists for more than 72 hours.
- If swallowed, immediately call Poison Control Center or doctor.

Keep out of reach of children.

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Directions

- Wet hands and forearms. Apply a palmful to hands and scrub hands and forearms thoroughly for 15 seconds or more. Rinse and repeat.

Other Inforamtion

- Store at 20-25°C (68-77°F)

Inactive Ingredients

Water, Alcohol Denat., Lauramine Oxide, Cocamidopropyl Betaine, Cetrimonium Chloride, Butylene Glycol, Citric Acid, Zinc Gluconate, PPG-2 Hydroxyethyl Cocamide, Farnesol, Glycerin, Methylchloroisothiazolinone, Polyaminopropyl Biguanide, Fragrance, Polyquaternium-10.

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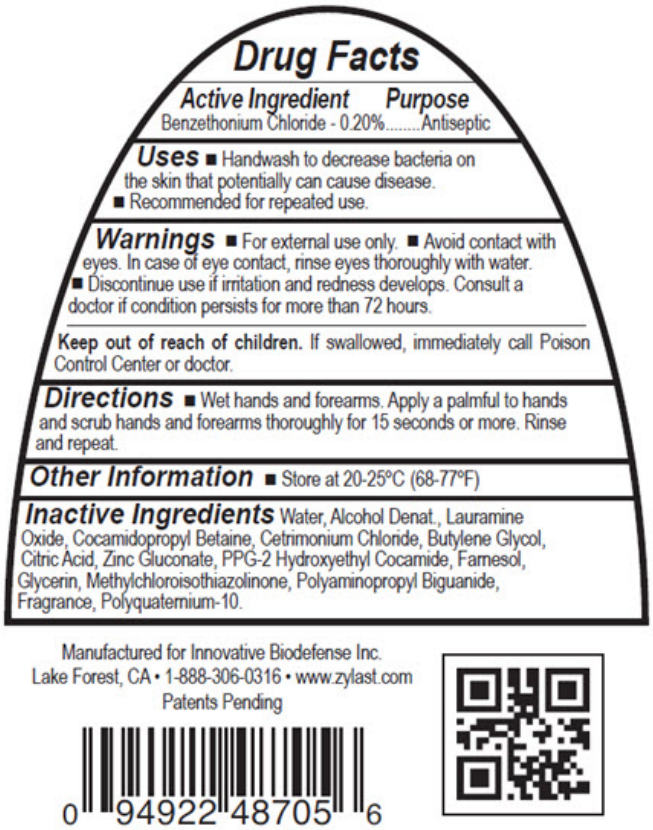
Package/Label Principal Display Panel

NDC 57702-487-05
Zylast XP
Extended Protection
Broad Spectrum
Antimicrobial
Antiseptic
Foaming Soap
8.25 oz 244mL



Package/Label Principal Display Panel

NDC 57702-487-15
Zylast XP
Extended Protection
Broad Spectrum
Antimicrobial
Antiseptic
Foaming Soap
1000mL (33.8oz)

ZYLAST XP ANTISEPTIC FOAMING

benzethonium chloride soap

Product Information			
Product Type	HUMAN OTC DRUG LABEL	**Item Code (Source)**	NDC:57702-487
Route of Administration	TOPICAL	**DEA Schedule**	

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZETHONIUM CHLORIDE (BENZETHONIUM)	BENZETHONIUM CHLORIDE	2 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER	
LAURAMINE OXIDE	
COCAMIDOPROPYL BETAINE	
CETRIMONIUM CHLORIDE	
BUTYLENE GLYCOL	
CITRIC ACID MONOHYDRATE	
ZINC GLUCONATE	
PPG-2 HYDROXYETHYL COCAMIDE	
FARNESOL	
GLYCERIN	
METHYLCHLOROISOTHIAZOLINONE	
POLIHEXANIDE	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:57702-487-05	244 mL in 1 BOTTLE		
2	NDC:57702-487-15	1000 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	06/01/2011	

Labeler - Bocchi Laboratories Inc. (078376306)

Revised: 3/2014 Bocchi Laboratories Inc.