

HILLYARD AMB SO AP- chloroxylenol liquid
Hillyard GMP

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.

HILLYARD® AMB Soap

Active ingredient

Chloroxylenol 0.3%

Purpose

Antimicrobial

Use

- Handwash to help decrease bacteria on the skin
- Recommended for repeated use

Warnings

For external use only

When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash appears and lasts

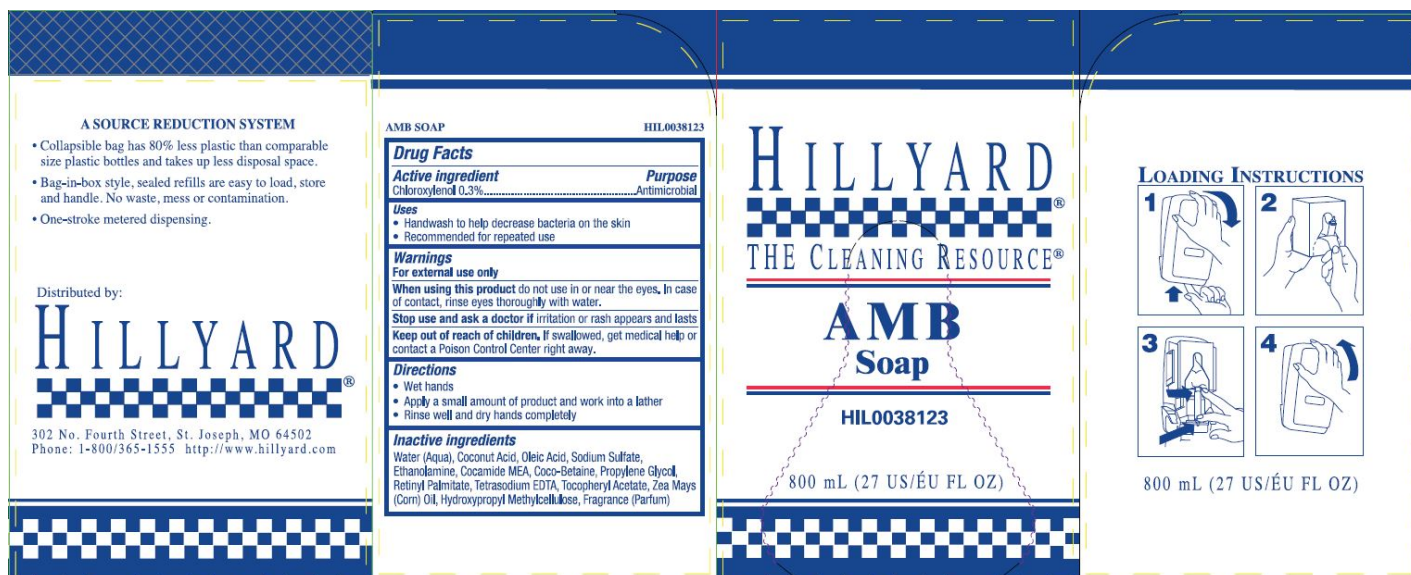
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Wet hands
- Apply a small amount of product and work into a lather
- Rinse well and dry hands completely

Inactive ingredients

Water (Aqua), Coconut Acid, Oleic Acid, Sodium Sulfate, Ethanolamine, Cocamide MEA, Coco-Betaine, Propylene Glycol, Retinyl Palmitate, Tetrasodium EDTA, Tocopheryl Acetate, Zea Mays (Corn) Oil, Hydroxypropyl Methylcellulose, Fragrance (Parfum)



HILLYARD AMB SO AP

chloroxylenol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76402-050
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Chloroxylenol (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	Chloroxylenol	0.003 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Coconut Acid (UNII: 40U37V505D)	
Oleic Acid (UNII: 2UMI9U37CP)	
Sodium Sulfate (UNII: 0YPR65R21J)	
MONOETHANOLAMINE (UNII: 5KV86114PT)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
Coco-Betaine (UNII: 03DH2IZ3FY)	
Propylene Glycol (UNII: 6DC9Q167V3)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
EDETATE SODIUM (UNII: MP1J8420LU)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CORN OIL (UNII: 8470G57WFM)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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1	NDC:76402-050-27	800 mL in 1 PACKAGE; Type 0: Not a Combination Product	04/30/2003	
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Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/30/2003	

Labeler - Hillyard GMP (969081483)

Revised: 8/2017

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