

CORALITE ULTRA ANTIBACTERIAL DEODORANT - triclocarban soap

United Exchange Corp

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.

Drug Facts

Active ingredient	Purpose
Triclocarban 0.3%.....	Antibacterial

Uses

- for washing to decrease bacteria on the skin

Warnings For external use only.

When using this product avoid contact with eyes. In case of eye contact, flush with water.

Stop use and ask a doctor if irritation and redness develops.

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

Directions

- wet bar with water
- lather vigorously and wash skin
- rinse and dry thoroughly

Inactive ingredients soap (sodium tallowate, sodium cocoate, sodium palmate types), water, glycerin, fragrance, sodium chloride, palm acid or tallowate acid, PEG-6 methyl ether, sorbitol, tetrasodium etidronate, BHT, pentasodium pentetate, FD&C yellow no. 5, D&C yellow no. 8, FD&C red no. 4

Distributed By:

United Exchange Corp.

17211 Valley View Ave.

Cerritos, CA 90703 USA

Made in Korea



CORALITE ULTRA ANTIBACTERIAL DEODORANT

triclocarban soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65923-807
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TRICLO CARBAN (UNII: BGG1Y1ED0 Y) (TRICLOCARBAN - UNII:BGG1Y1ED0 Y)	TRICLOCARBAN	0.3 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
SODIUM COCOATE (UNII: R1TQH25F4I)	
SODIUM PALMATE (UNII: S0A6004K3Z)	
WATER (UNII: 059QF0K00R)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
PALM ACID (UNII: B6G0Y5Z616)	
PEG-6 METHYL ETHER (UNII: WXH089JZ5E)	
SORBITOL (UNII: 506T60A25R)	
PENTASODIUM PENTETATE (UNII: 961TOZ5L7T)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
D&C YELLOW NO. 8 (UNII: 93X55PE38X)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65923-807-35	100 g in 1 PACKAGE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/30/2014	

Labeler - United Exchange Corp (840130579)

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