NOXZEMA BACTERIA FIGHTING CLEANSER- triclosan soap CONOPCO Inc. d/b/a Unilever

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.

NOXZEMA®

Drug Facts

Active ingredient

Triclosan 0.3%

Purpose

Antibacterial

Uses

For cleansing to decrease bacteria on skin

Warnings

• **For external use only.** Do not use on infants under 6 months of age.

When using this product

- Avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water.
- Discontinue use if signs of irritation or rash appear.

Stop use and ask a doctor if irritation or rash persists or is excessive.

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

Directions

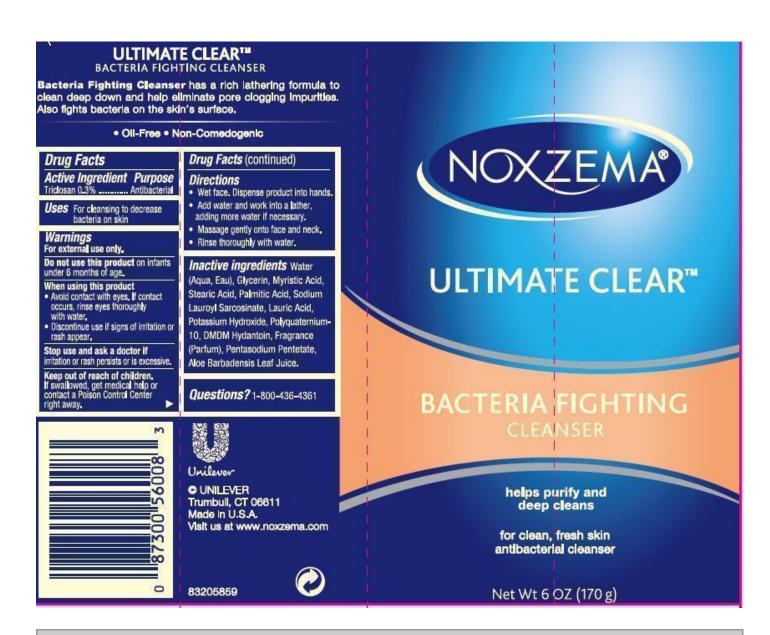
- Wet face. Dispense product into hands.
- Add water and work into a lather, adding more water if necessary.
- Massage gently onto face and neck.
- Rinse thoroughly with water.

Inactive ingredients

Water (Aqua, Eau), Glycerin, Myristic Acid, Stearic Acid, Palmitic Acid, Sodium Lauroyl Sarcosinate, Lauric Acid, Potassium Hydroxide, Polyquaternium-10, DMDM Hydantoin, Fragrance (Parfum), Pentasodium Pentetate, Aloe Barbadensis Leaf Juice.

Questions?

1-800-436-4361



NOXZEMA BACTERIA FIGHTING CLEANSER

triclosan soap

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64942-1264
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
Triclosan (UNII: 4NM5039 Y5X) (Triclosan - UNII:4NM5039 Y5X)	Triclosan	0.003 g in 1 g

Inactive Ingredients			
Ingredient Name	Strength		
Water (UNII: 059QF0KO0R)			
Aloe Vera Leaf (UNII: ZY8 1Z8 3H0 X)			
Glycerin (UNII: PDC6A3C0OX)			

Iodopropynyl Butylcarbamate (UNII: 603P14DHEB)	
Palmitic Acid (UNII: 2V16EO95H1)	
Myristic Acid (UNII: 013V7S25AW)	
Stearic Acid (UNII: 4ELV7Z65AP)	
Lauric Acid (UNII: 1160N9NU9U)	
Potassium Hydroxide (UNII: WZH3C48M4T)	

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:64942-1264-1	170 g in 1 PACKAGE		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	10/02/2012	

Labeler - CONOPCO Inc. d/b/a Unilever (001375088)

Establishment			
Name	Address	ID/FEI	Business Operations
Alberto-Culver USA Inc.		021679448	MANUFACTURE(64942-1264)

Establishment			
Name	Address	ID/FEI	Business Operations
Cosmetic Laboratories of America LLC		013696501	manufacture(64942-1264)

Revised: 10/2012 CONOPCO Inc. d/b/a Unilever