ANTIBACTERIAL LIQUID HAND - triclos an liquid Vi Jon

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

Triclosan 0.115%

Purpose

Antibacterial

Uses for handwashing to decrease bacteria on the skin

Warnings

For external use only.

When using this product • do not get into eyes. If contact occurs, rinse eyes thoroughly with water.

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

Directions •Wet hands •Apply palmful to hands •Scrub thoroughly •Rinse

Inactive ingredients water, sodium laureth sulfate, sodium lauryl sulfate, cocamidopropyl betaine, sodium chloride, decyl glucoside, fragrance, DMDM hydantoin, PEG-120 methyl glucose dioleate, cocamide MEA, citric acid, tetrasoidum EDTA, polyquaternium-7, PEG-7 glyceryl cocoate, D+C red no. 33, FD+C, blue no. 1

Distributed by: Vi Jon 8515 Page Ave., St. Louis, MO. 63114

This product is not manufactured or distributed by Colgate-Palmolive Company, distributor of Softsoap Aquarium Series Antibacterial Soap

Swan
CLEAR ANTIBACTERIAL
HAND SOAP
Compare to Softsoap
Aquarium Series Antibacterial Soap
11.25 FL OZ (332 mL)
Compare to Softsoap
Aquarium Series Antibacterial Soap





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11.25 FL OZ (332 mL)

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ANTIBACTERIAL LIQUID HAND

triclosan liquid

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:11344-541

Route of Administration TOPICAL

Active Ingredient/Active Moiety				
l	Ingredient Name	Basis of Strength	Strength	
l	TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X)	TRICLOSAN	.115 kg in 100 L	

Inactive Ingredients				
Ingredient Name	Strength			
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)				
SODIUM LAURYL SULFATE (UNII: 368GB5141J)				
COCAMIDO PRO PYL BETAINE (UNII: 50 CF3 O 11 KX)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
DMDM HYDANTO IN (UNII: BYR0546 TOW)				
PEG-120 METHYL GLUCO SE DIOLEATE (UNII: YM0 K64F20 V)				

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:11344-541-93	1.18 L in 1 BOTTLE, PLASTIC			
2	NDC:11344-541-96	.221 L in 1 BOTTLE, PLASTIC			
3	NDC:11344-541-81	.332 L in 1 BOTTLE, PLASTIC			

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333A	07/30/2009			

Labeler - Vi Jon (150931459)

Registrant - Vi Jon (150931459)

Establishment					
Name	Address	ID/FEI	Business Operations		
Vi Jon		150931459	manufacture		

Revised: 11/2010 Vi Jon