ANTIBACTERIAL LIQUID HAND - triclos an liquid OnPoint, 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.

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 OnPoint, Inc 2 Paragon Drive, Montvale, NJ 07645

MARKET SPA ANTIBACTERIAL LIQUID HAND SOAP CLEAR FORMULA 7.5 FL OZ (221 mL)



ANTIBACTERIAL LIQUID HAND

triclosan liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:51143-541

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TRICLOSAN (LINII) ANMED 20 VEY) (TRICLOSAN LINII) ANMED 20 VEY)	TRICLOSAN	115 kg in 100 I

.115 kg in 100 L

Inactive Ingredients

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Ingredient Name	Strength				
SO DIUM LAURETH SULFATE (UNII: BPV390 UAP0)					
SODIUM LAURYL SULFATE (UNII: 368GB5141J)					
COCAMIDO PRO PYL BETAINE (UNII: 50 CF3 O 11 KX)					
SO DIUM CHLO RIDE (UNII: 451W47IQ8X)					
DMDM HVDANTO IN (LINII) DVD0546 TOWO					

DMDM HYDANTO IN (UNII: BYR0546 TOW)

PEG-120 METHYL GLUCO SE DIO LEATE (UNII: YM0 K6 4F20 V)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51143-541-93	1.18 L in 1 BOTTLE, PLASTIC		
2	NDC:51143-541-96	.221 L in 1 BOTTLE, PLASTIC		
3	NDC:51143-541-97	1.89 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 - OnPoint, Inc (001367366)

Registrant - Vi Jon (150931459)

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Name	Address	ID/FEI	Business Operations		
Vi Jon		150931459	manufacture		

Revised: 11/2010 OnPoint, Inc