FOAMING HAND WASH ANTIBACTERIAL- triclos an liquid Target Corporation

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 Ingredients

Triclosan 0.6%

Purpose

Antibacterial

Use

Uses for hand washing to decrease bacteria on the skin

Warnings

Warnings

For external use only-hands only

When using this product

When using this product

• avoid contact with the eyes. If contact occurs, rinse eyes with water.

Stop use and ask a doctor

Stop use and ask a doctor if irritation or redness develops

Keep out or reach of children

Keep out or freach of children except under adult supervision. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Directions

- wet hands
- apply palmful to hands
- scrub thoroughly
- rinse thoroughly

Inactive ingredients

Inactive ingredients water, sodium laureth sulfate, glycerin, cocamidopropyl betaine, ammonium xylenesulfonate, fragrance, disodium phosphate, citric acid, benzyl alcohol, red 4, yellow 5

Questions

Questions? Call 1-800-910-6874

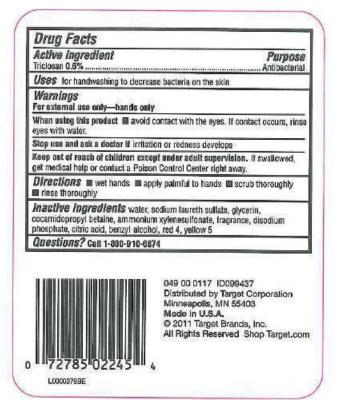
Adverse reactions

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Minneapolis, MN 55403
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Principal display panel

foaming hand wash
antibacterial
gently cleans hand
refill for foaming soap dispensers
Loaded Lather
up + up
50 FL OZ (1.4 L)





FOAMING HAND WASH ANTIBACTERIAL

triclosan liquid

Product Information			
Product Type	HUMAN OTC DRUG LABEL	Item Code (Source)	NDC:11673-383
Route of Administration	TOPICAL	DEA Schedule	

l	Active Ingredient/Active Moiety			
l	Ingredient Name	Basis of Strength	Strength	
ı	TRICLOSAN (TRICLOSAN)	TRICLOSAN	6 mg in 1 mL	

Inactive Ingredients			
Ingredient Name	Strength		
WATER			
SO DIUM LAURETH SULFATE			
GLYCERIN			
CO CAMIDO PRO PYL BETAINE			
AMMO NIUM XYLENES ULFO NATE			
SO DIUM PHO SPHATE, DIBASIC, ANHYDRO US			
CITRIC ACID MONOHYDRATE			
BENZYL ALCOHOL			
FD&C RED NO. 4			
FD&C YELLOW NO. 5			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:11673-383-03	1478 mL in 1 BOTTLE, PUMP			
2	NDC:11673-383-96	.221 mL in 1 PACKAGE			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	03/19/1993		

Labeler - Target Corporation (006961700)

Registrant - Vi-Jon (790752542)

Establishment				
Name	Address	ID/FEI	Business Operations	
Vi-Jon		088520668	manufacture(11673-383)	

Revised: 10/2012 Target Corporation