KEYSTONE - triclosan solution
Ecolab Inc.

Drug Facts

Active ingredient

Triclosan 0.3%

Purpose

Antiseptic handwash

Uses

- for handwashing to decrease bacteria on the skin

Warnings
For external use only

Do not use

- in eyes

When using this product

- if in eyes, rinse promptly and thoroughly with water
- discontinue use if irritation and redness develop

Stop use and ask a doctor if

- skin irritation or redness occurs for more than 72 hours

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

Directions

- wet skins and apply a small amount on hands an forearms
- scrub well, rinse thoroughly and dry

Other information

- for additional information see, Safety Data Sheet (SDS)
- for emergency medical information in USA and Canada, call 1.800.328.0026

Inactive ingredients water (aqua), potassium cocoate, alcohol, sodium laureth sulfate, hexylene glycol, boric acid, cocamidopropyl PG-dimonium chloride phosphate, tetrasodium EDTA, cocamine oxide, fragrance, PEG-75 lanolin, methylparaben, propylparaben, aloe barbadensis leaf juice, FD&C blue 1

Questions? call **1.800.35.CLEAN (352.5326)**

Principal display panel and representative label

KEYSTONE

ANTIBACTERIAL FOAM HANDSOAP

Triclosan 0.3%

SYSCO 0281531

6100517 ECOLAB

Net Contents: 1200 mL (40.6 US fl oz)

750767/5401/0420

Distributed by

Sysco Corporation

Manufactured by

Ecolab · 1 Ecolab Place · St. Paul MN 55102 USA ·

tel: 1 800 35 CLEAN (352 5326)

Made in U.S.A.

Keystone Antibacterial Foam Handsoap

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750768/540V/0420

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This product may be patented: www.ecolab.com/patents

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KEYSTONE
triclosan solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47593-341
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X)	TRICLOSAN	0.3 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
POTASSIUM COCOATE (UNII: F8U72V8ZXP)	
ALCOHOL (UNII: 3K9958V90M)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
BORIC ACID (UNII: R57ZHV85D4)	
COCAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
EDETATE SODIUM (UNII: MP1J8420LU)	
COCAMINE OXIDE (UNII: QWA2IZI6FI)	
PEG-75 LANOLIN (UNII: 09179OX7TB)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47593-341-41	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/26/2000	05/03/2024
2	NDC:47593-341-56	1200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/26/2000	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	07/26/2000	

Labeler - Ecolab Inc. (006154611)

Revised: 3/2024 Ecolab Inc.