

BACTI-STAT AE- triclosan solution
Ecolab 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, 0.3%

Purpose

Healthcare personnel handwash

Uses

- healthcare personnel handwash

Warnings

- **For external use only**

Do not use

- in eyes

When using this product

- discontinue use if irritation and redness develop

Stop use and ask a doctor if

- skin irritation or redness persists 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 skin and spread a small amount on hands and forearms
- scrub well, rinse thoroughly and dry

Other information

- for additional information, see Material Safety Data Sheet (MSDS)
- for emergency medical information in USA and Canada, call 1.800.328.0026
- for emergency medical information worldwide, call 1 651 222 5352 (in the USA)

☐ **Inactive ingredients** water, potassium cocoate, SD alcohol 40-B, glycerin, cocamidopropyl PG-dimonium chloride phosphate, potassium stearate, tetrasodium EDTA, hydroxyethylcellulose, fragrance, cocamine oxide, methylparaben, tocopheryl acetate, citric acid, isopropyl alcohol, propylparaben, aloe barbadensis leaf juice, FD&C blue 1, FD&C yellow 5

☐ **Questions?** ☐ call **1.866.781.8787**

Principal Display Panel and Representative Label

ECOLAB

NDC 47593-170-31

BACTI-STAT AE®

Antimicrobial Hand Soap

Formulated with Aloe Vera & Vitamin E

to promote skin health

Healthcare personnel handwash

Active Ingredient: 0.3% Triclosan

6060116

Net Contents: 540 mL (18 fl oz)

Ecolab · 370 Wabasha Street N ·
St Paul MN 55102-1390 USA

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Made in United States

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SDS-IN-1633

ECOLAB®



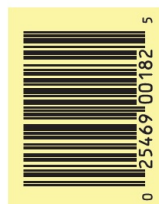
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Questions? call 1 866 781 8787	

BACTI-STAT AE

triclosan solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47593-170
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Triclosan (UNII: 4NM5039Y5X) (Triclosan - UNII:4NM5039Y5X)	Triclosan	3.06 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
POTASSIUM COCOATE (UNII: F8U72V8ZXP)	
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
COCAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
POTASSIUM STEARATE (UNII: 17V812XK50)	
EDETATE SODIUM (UNII: MP1J8420LU)	
HYDROXYETHYL CELLULOSE (2000 CPS AT 1%) (UNII: S38J6RZN16)	
COCAMINE OXIDE (UNII: QWA2IZI6FI)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47593-170-31	540 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/21/2012	
2	NDC:47593-170-32	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/21/2012	
3	NDC:47593-170-26	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/21/2012	
4	NDC:47593-170-56	1200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/12/2009	04/27/2016
5	NDC:47593-170-41	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/22/2013	04/27/2016
6	NDC:47593-170-59	1250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/22/2013	04/27/2016

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/14/1997	

Labeler - Ecolab Inc. (006154611)