DIGISAN E- benzalkonium chloride 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

Benzalkonium chloride, 0.1%

Purpose

Antiseptic handwash

Uses

- for handwashing to decrease bacteria on the skin

Warnings

- **For external use only**
- **Flammable, keep away from fire or flame, heat sparks and sources of static discharge**

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 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 hands and remove soil
- dispense palmful
- spread to cover hands, rub in well
- air dry, do not rinse or towel dry

Other Information

- For additional information, see Safety Data Sheet (SDS)
- For emergency medical information in USA and Canada, call 1-800-328-0026
- For emergency medical information worldwide, call 1-651-222-5352 (in USA)

Inactive ingredients:

water (aqua), isopropyl alcohol, propylene glycol, FD&C red 40, F&DC blue 1

Questions?

Call 1-800-35-CLEAN (352-5326)

Principal Display Panel/Representative Label

DIN 02242842

ECOLAB

23674

6123674

DigiSan™

E Foam Hand Sanitizer

750 mL (25 US FL OZ)

Active Ingredient:

Benzalkonium chloride 0.1%

706304/5404/0117

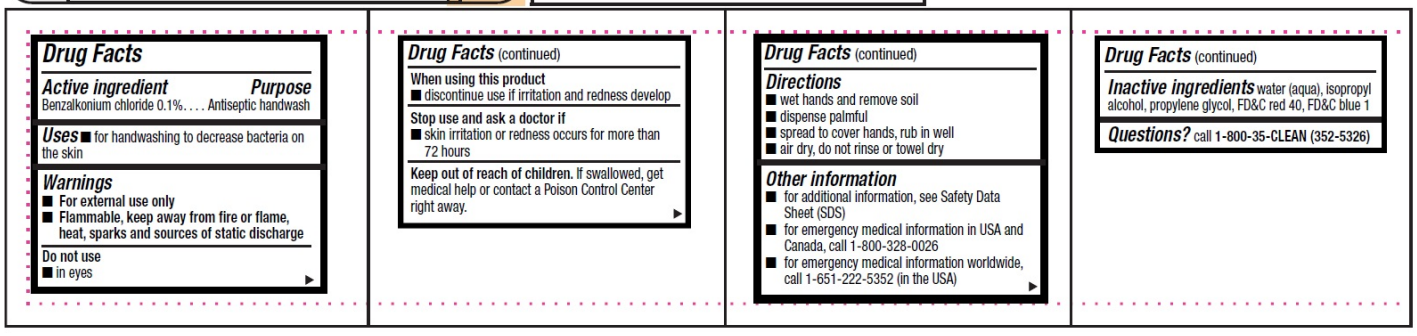
This product may be patented | Ce produit peut être breveté | Este producto puede ser patentado: www.ecolab.com/patents

Ecolab · 1 Ecolab Place · St. Paul MN 55102 USA · tel: 1 800 35 CLEAN (352 5326)

5105 Tomken Road · Mississauga ON L4W 2X5 Canada

Made in U.S.A. | Fabriqué aux É-U | Hecho en EE.UU.

www.ecolab.com · 706304/5404/0117



DIGISAN E

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47593-404
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47593-404-30	207 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/04/2003	03/13/2017
2	NDC:47593-404-41	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/04/2003	
3	NDC:47593-404-59	1250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/04/2003	03/13/2017

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	11/04/2003	

Labeler - Ecolab Inc. (006154611)

Revised: 3/2017 Ecolab Inc.