

ECOLAB FOAMING ANTIBACTERIAL HANDSOAP- chloroxylenol solution
Ecolab Inc.

Drug Facts

Active ingredient

Chloroxylenol 0.5%

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 hands and forearms. • Apply palmful to hands and forearms.
- Scrub thoroughly. • Rinse and dry.

Other information

- for additional information, see Safety Data Sheet (SDS)
- **Medical Emergency:** (877) 231-2615 or call collect 0 (952) 853-1713

Inactive Ingredients

water (aqua), potassium cocoate, hexylene glycol, sodium sulfate, tetrasodium EDTA, sodium lauryl sulfate, glycerin, citric acid, coco-glucoside, glyceryl oleate, fragrance,

methylchloroisothiazolinone, CI 19140 (FD&C Yellow No. 5), methylisothiazolinone, CI 14700 (FD&C Red. No. 4)

Questions? Call 1-800-529-5458

Representative Label and Principal Display Panel

ECOLAB®

Peel Here For Drug Facts

NDC 63146-124-10

Foaming Antibacterial

Hand Soap

Jabón de manos

antibacterial en espuma

Foaming Antibacterial Hand Soap is a rich lather hand soap with a balanced blend of cleaning agents and skin protecting moisturizers.

KEEP OUT OF REACH OF CHILDREN • FOR INSTITUTIONAL USE ONLY
Chloroxylenol 0.5%

To obtain Spanish Instructions, see outer carton.

Para obtener las instrucciones en español, véase la caja exterior.

Net contents: 42 US fl oz (1250 ml)

Distributed by:

Kay Chemical Company · 8300 Capital Drive

Greensboro, NC 27409-9790 USA

Customer Service: (800) 529-5458

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FRSUSA 764645/8001/0221

Drug Facts (continued)

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Drug Facts (continued)

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ECOLAB FOAMING ANTIBACTERIAL HANDSOAP

chloroxylenol solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47593-678
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
POTASSIUM COCOATE (UNII: F8U72V8ZXP)	

HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
SODIUM SULFATE ANHYDROUS (UNII: 36KCS0R750)	
EDETATE SODIUM (UNII: MP1J8420LU)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
GLYCERIN (UNII: PDC6A3C0OX)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
COCO-GLUCOSIDE (UNII: ICS790225B)	
GLYCERYL OLEATE (UNII: 4PC054V79P)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47593-678-59	1250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/18/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	12/18/2025	

Labeler - Ecolab Inc. (006154611)

Revised: 12/2025

Ecolab Inc.