

KAY FOAMING ANTIBACTERIAL HANDSOAP- chloroxyenol 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 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

- EMERGENCY HEALTH INFORMATION: 1 800 328 0026 (number is in the US).
- If located outside the United States and Canada, call collect 1 651 222 5352

Inactive ingredients water (aqua), potassium cocoate, hexylene glycol, sodium sulfate, tetrasodium EDTA, sodium lauryl sulfate, glycerin, sodium citrate, glyceryl oleate, fragrance, sodium glycolate, caprylyl/capryl glucoside, lauryl glucoside, methylchloroisothiazolinone, yellow 5, methylisothiazolinone, red 4

Questions?

Questions? Call **1-800-529-5458**

Representative label and Principal Display Panel

ECOLAB

Kay Foaming

Antibacterial Handsoap

Active Ingredient: Chloroxylenol 0.5%

Instant foam lathers fast • Mild, pH-balanced formula

Fresh, clean fragrance

KAY® Foaming Antibacterial Handsoap 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

Manufactured by: Ecolab Inc

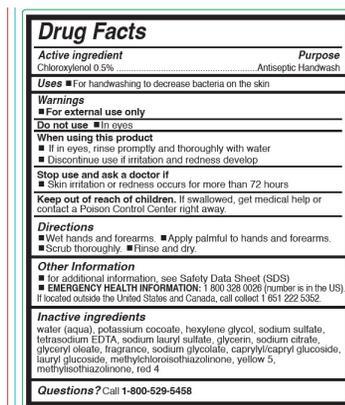
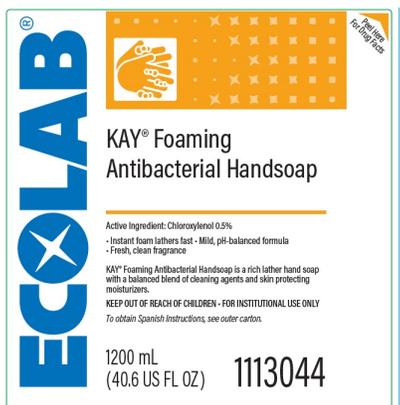
1 Ecolab Place · St. Paul, MN 55102 USA

Customer Service: (800) 529-5458

©2025 Kay Chemical Company

All rights reserved | Made in USA

KUSA 785111/8000/0925



KAY FOAMING ANTIBACTERIAL HANDSOAP

chloroxylenol solution

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:47593-661

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)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
GLYCERYL OLEATE (UNII: 4PC054V79P)	
SODIUM GLYCOLATE (UNII: B75E535IMI)	
CAPRYLYL/CAPRYL GLUCOSIDE (UNII: E00JL9G9K0)	
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
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:47593-661-70	1200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/04/2027	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	10/04/2025	

Labeler - Ecolab Inc. (006154611)

Revised: 12/2025

Ecolab Inc.