ECOLAB FOAMING AB HS- chloroxylenol liquid
Kay Chemical Company

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

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

Other information

- for additional infomration see Safety Data Sheet (SDS)
- EMERGENCY HEALTH INFORMATION: 1 877 231 2615. If located outside the United States and Canada, call collect 952 853 1713 (number is in the US).

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, magnesium nitrate, sodium hydroxide, methylchloroisothiazolinone, magnesium chloride, yellow 5, methylisothiazolinone, red 4

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

Principal display panel and representative label

ECOLAB

NDC 63146-319-16

Foaming AB Hand Soap

Active Ingredient: Chloroxylenol 0.5%

1200 mL

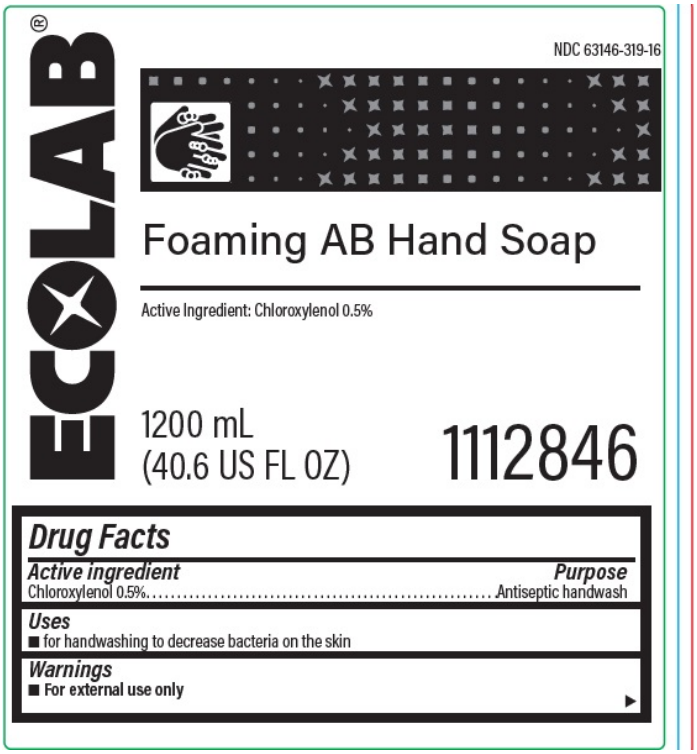
(40.6 US FL OZ) 1112846

Kay Chemical Company · 8300 Capital Drive

Greensboro, NC 27409-9790 USA

Customer Service: (800) 529-5458

KUSA · 782978/8000/0224



ECOLAB FOAMING AB HS

chloroxylenol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63146-319
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 (UNII: 0YPR65R21J)	
EDETATE SODIUM (UNII: MP1J8420LU)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
GLYCERYL OLEATE (UNII: 4PC054V79P)	
SODIUM GLYCOLATE (UNII: B75E535IMI)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
MAGNESIUM NITRATE (UNII: 77CBG3UN78)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
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:63146-319-16	1200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/19/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	06/19/2024	

Labeler - Kay Chemical Company (003237021)

Revised: 6/2024

Kay Chemical Company