KAY MCD- chloroxylenol solution
Kay Chemical Company

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

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 and 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 5 ml (teaspoonful) or palmful to hands and forearms.
- Scrub thoroughly for 20 seconds.
- Rinse and repeat.

Other information

- For additional information, see Material Safety Data Sheet (MSDS)

- For emergency medical information in USA call (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, hydroxyethylcellulose, glycerine, coco glucoside, glyceryl oleate, citric acid, fragrance, methylchloroisothiazolinone, methylisothiazolinone, FD&C Red 40, FD&C Yellow 5, D&C Red 33.

Questions?

Call 1-800-529-5458

Principal Display Panel - Representative Label

NDC No.: 63146-102-01

McD

Anti-Microbial

Handwash (AMH)

FOR INSTITUTIONAL USE ONLY

(Chloroxylenol 0.5%)

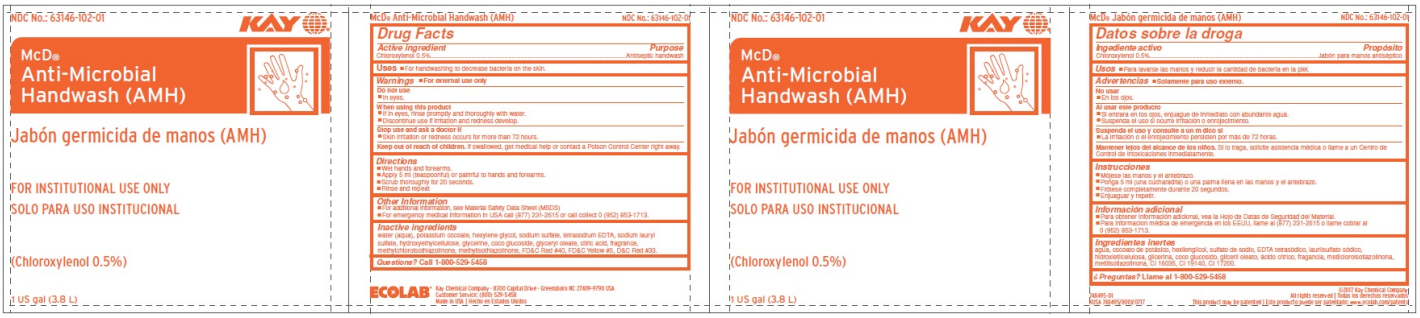
1 US gal (3.8L)

ECOLAB

KAY CHEMICAL COMPANY - 8300 Capital Drive - Greensboro, NC 27409-9790 USA

Customer Service: (800) 529-5458

Made in USA



KAY MCD

chloroxylenol solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63146-102

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.5 mg in 100 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)	
HYDROXYETHYL CELLULOSE (3000 MPA.S AT 1%) (UNII: 7Q6P4JN1QT)	
GLYCERIN (UNII: PDC6A3C0OX)	
COCO GLUCOSIDE (UNII: ICS790225B)	
GLYCERYL OLEATE (UNII: 4PC054V79P)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63146-102-01	3800 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/13/2005	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	10/13/2005	

Labeler - Kay Chemical Company (003237021)

Revised: 7/2022 Kay Chemical Company