

**SYSCO RELIANCE- chloroxylenol solution**  
**Ecolab Inc.**

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**Drug Facts**

**Active ingredient**

Chloroxylenol, 0.5%

**Purpose**

Antiseptic handwash

**Uses**

- for handwashing to decrease bacteria on the skin
- recommended for repeated use

**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
- dispense a palmful of product to hands
- scrub hands and forearms for 20 seconds
- rinse thoroughly and dry

**Other information**

- for additional information, see Safety Data Sheet (SDS)
- for emergency medical information in USA and Canada, call 1.800.328.0026

**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 (FDC Yellow No. 5), methylisothiazolinone, CI 14700 (FDC Red No. 4)

Questions? call 1.800.35.CLEAN (352.5326)

**Principal display panel and representative label**

Sysco Reorder #

Reliance 0279754

AB FOAM HAND SOAP

ACTIVE INGREDIENT: CHLOROXYLENOL 0.5%

DISTRIBUTED BY SYSCO CORPORATION

HOUSTON, TEXAS 77077

001123 6100592

751544/5404/1122

NET CONTENTS: 25.4 US FL OZ (750 mL)

**Other information**

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**0279754**

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**Drug Facts** (continued)

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**SYSCO RELIANCE**

chloroxylenol solution

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:47593-477
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>CHLOROXYLENOL</b> (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.5 mg in 100 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>POTASSIUM COCOATE</b> (UNII: F8U72V8ZXP)	
<b>HEXYLENE GLYCOL</b> (UNII: KEH0A3F75J)	
<b>SODIUM SULFATE</b> (UNII: 0YPR65R21J)	
<b>EDETATE SODIUM</b> (UNII: MP1J8420LU)	
<b>SODIUM LAURYL SULFATE</b> (UNII: 368GB5141J)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	
<b>COCO GLUCOSIDE</b> (UNII: ICS790225B)	
<b>GLYCERYL OLEATE</b> (UNII: 4PC054V79P)	
<b>METHYLCHLOROISOTHIAZOLINONE</b> (UNII: DEL7T5QRPN)	
<b>FD&amp;C YELLOW NO. 5</b> (UNII: I753WB2F1M)	
<b>METHYLISOTHIAZOLINONE</b> (UNII: 229D0E1QFA)	
<b>FD&amp;C RED NO. 4</b> (UNII: X3W0AM1JLX)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47593-477-41	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/05/2011	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	04/05/2011	

**Labeler** - Ecolab Inc. (006154611)

Revised: 12/2025

Ecolab Inc.