

**DYNAREX GREEN- chloroxylenol liquid**  
**Dynarex Corporation**

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**1438 Dynarex Antibacterial Hand Soap NDC 67777-318-01**

**ACTIVE INGREDIENT**

Active Ingredient	Purpose
Chloroxylenol 0.15%	Antiseptic

**WARNINGS**

- **For external use only**
- **When using this product** avoid contact with eyes. In case of contact, flush with water.

**PURPOSE**

An antiseptic handwash.

**Stop use and ask a doctor if**

**Stop use and ask a doctor if** irritation and redness develop.

**INDICATIONS & USAGE**

Cleaning and antiseptic cleansing of hands and skin.

**DOSAGE & ADMINISTRATION**

- Wet hands and forearms. Apply 5 milliliters (teaspoon) or palmful to hands and forearms. Scrub thoroughly for 1 minute. Rinse and repeat.

**KEEP OUT OF REACH OF CHILDREN**

**KEEP OUT OF REACH OF CHILDREN,** if swallowed get medical help or contact a Poison Control Center right away.

**INACTIVE INGREDIENTS**

Inactive ingredients: Citric acid, Cocamidopropyl betaine, Cocmide DEA, D&C Red #33, Disodium EDTA, Frgarance, Glycerin, Sodium chloride, Sodium Lauryl Sulphate, Sodium o-phenylphenate, water.

**PRINCIPAL DISPLAY PANEL**

Dynarex Antibacterial Soap

1438 Soap.jpg





## DYNAREX GREEN

chloroxylenol liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:67777-318
<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	0.15 g in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>COCAMIDOPROPYL BETAINE</b> (UNII: 5OCF3O11KX)	
<b>COCO DIETHANOLAMIDE</b> (UNII: 92005F972D)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>SODIUM LAURETH SULFATE</b> (UNII: BPV390UAP0)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM O-PHENYLPHENATE</b> (UNII: KJV9K7N7UI)	
<b>D&amp;C RED NO. 33</b> (UNII: 9DBA0SBB0L)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-318-01	221 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2017	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC Monograph Drug M003

03/13/2017

**Labeler** - Dynarex Corporation (008124539)

**Registrant** - Dynarex Corporation (008124539)

Revised: 11/2024

Dynarex Corporation