DYNAREX GREEN- chloroxylenol liquid
Dynarex Corporation

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.

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			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-318
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.15 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
COCO DIETHANOLAMIDE (UNII: 92005F972D)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
WATER (UNII: 059QF0KO0R)	
SODIUM O-PHENYLPHENATE (UNII: KFV9K7N7UI)	
D&C RED NO. 33 (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 not final	part333A	03/13/2017	

Labeler - Dynarex Corporation (008124539)

Registrant - Dynarex Corporation (008124539)

Revised: 12/2021

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