DYNAREX GREEN- alcohol 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.

1362 Dynarex Green Soap NDC 67777-308-01
1363 Dynarex Green Soap NDC 67777-308-02

ACTIVE INGREDIENT

Active Ingredient	Purpose
Alcohol, 30% v/v	Antiseptic

WARNINGS

- **For external use only**
- Flammable, keep away from fire or flame
- Avoid contact with eyes and mucousal membranes

DO NOT USE

Do not use with electrocautery procedures

PURPOSE

An antiseptic wash.

INDICATIONS & USAGE

Cleaning and antiseptic cleansing of skin.

DOSAGE & ADMINISTRATION

Mix with water, lather and rinse off.

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.

OTHER INFORMATION

- Store at room temperature: 15 º - 30 º C (59 º - 86 º F)
- Avoid excessive heat

INACTIVE INGREDIENTS

Inactive ingredients: Coconut acid, fragrance, glycerin, oleic acid, potassium hydroxide, water.

Label

Label

Reorder **No. 1362**

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LOT

dynarex

Tincture of Green Soap U.S.P.

Drug Facts

Active ingredient — **Purpose**
Alcohol 30% v/v ..... Antiseptic

Uses
- Cleaning and antiseptic cleansing of skin

Warnings
- **For External Use Only**
- **Flammable, keep away from fire or flame**

When using this product avoid contact with eyes or mucous membranes

Stop use and ask a doctor if
- Irritation and redness develop
- If condition persists for more than 72 hours

Keep out of reach of children. If swallowed, get medical help or contact Poison Control Center right away.

Do not use with electrocautery procedures

Directions
- Mix with water, lather and rinse

Other information
- Store at room temperature between 15°-30°C (59°-86°F)

Inactive ingredients
Fragrance, Glycerin, Oleic Acid, Potassium Hydroxide, Sunflower Oil, Water

R201202

SYMBOL GLOSSARY
For an explanation of symbols used in Dynarex packaging, visit dynarex.com/symbols.php

Manufactured for:
Dynarex Corporation
10 Glenshaw Street
Orangeburg, NY 10962
USA • www.dynarex.com

Made in India

16 fl. oz.
(473 ml)

DYNAREX GREEN
alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-308
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.3 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
OLEIC ACID (UNII: 2UMI9U37CP)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	

WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-308-11	12 in 1 CASE	07/01/2014	
1	NDC:67777-308-01	473 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:67777-308-12	4 in 1 CASE	07/01/2014	
2	NDC:67777-308-02	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:67777-308-03	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/01/2014	

Labeler - Dynarex Corporation (008124539)

Registrant - Dynarex Corporation (008124539)

Revised: 11/2022 Dynarex Corporation