DYNAREX HYDROGEL- glycerin gel
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.

1280 DynaGel Moisturizing Wound Hydrogel NDC 67777-233-01

Active Ingredient

Active Ingredient	Purpose
Glycerin 20.0%	Skin Protectant

Purpose

For dressing and management of stasis ulcers, pressure ulcers (stages I-IV) 1st and 2nd degree burns, cuts abrasions, skin irritations, post operative incisions, and skin conditions associated with peristomal care.

Warnings

- If condition worsens or does not improve within 10-14 days, consult a physician.
- Keep this and all medications out of the reach of children.
- Follow directions for use.

Dosage & Administration Dynarex Hydrogel

- Apply Wound Gel liberally to cover involved areas; apply as often as necessary.
- If gauze is used as a wound covering, moisten first.

Keep out of reach of children

- **KEEP THIS AND ALL MEDICATIONS OUT OF THE REACH OF CHILDREN**

Indications and Usage:

Use:

- For dressing and management of stasis ulcers,
- pressure ulcers (stages I-IV)
- 1st and 2nd degree burns,
- cuts abrasions,
- skin irritations,
- post operative incisions,
- and skin conditions associated with peristomal care.

Inactive Ingredient

Inactive ingredients: Allantoin, Aloe Vera Gel, Diazolidinyl Urea, Methyparaben, PEG-4 Olivate, PEG-60 Hydrogenated Castor Oil, Propylene Glycol, Propylparaben, Purified Water, Sodium Polyacrylate, Tetrasodium EDTA, Tocopherol Acetate.

We Care™ from dynarex

DYNAGEL

MOISTURIZING WOUND HYDROGEL

• *With Vitamin E and Aloe Vera*
• *Clean and Greaseless*

Net Wt. 3 oz.
(84.7 g)

Drug Facts

Active Ingredient — **Purpose**
Glycerin 20% ..... Skin Protectant

Use(s)
- Temporarily protects and helps relieve minor skin irritation and itching of minor cuts, scrapes, and burns
- Maintains a moist wound environment
- Ideal for management of stasis ulcers, pressure ulcers (stages 1-4), first and second-degree burns, cuts, post-operative incisions, and skin conditions associated with peristomal care

Warnings

For external use only

Do not use on ■ Deep or puncture wounds ■ Animal bites

When using this product ■ Avoid contact with eyes.

Stop use and ask doctor if ■ Condition worsens or does not improve within 10-14 days

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions
- Apply liberally as often as necessary to cover affected areas.
- If gauze is used as a wound covering, moisten first.
- Children under 6 months: Consult a doctor before use.

Other Information
- Store at room temperature 15°-30°C (59°-86°F)
- Tamper evident. Do not use if seal is damaged.

Inactive Ingredients
Allantoin, Aloe Vera Gel, Diazolidinyl Urea, Methylparaben, PEG-4 Olivate, PEG-60 Hydrogenated Castor Oil, Propylene Glycol, Propylparaben, Purified Water, Sodium Polyacrylate, Tetrasodium EDTA, Tocopherol Acetate

Questions?
1-888-DYNAREX Monday-Friday, 9AM-5PM EST

PATIENT NAME:

ROOM #

Reorder No. 1280

Manufactured for:
Dynarex Corporation
10 Glenshaw Street
Orangeburg, NY 10962
USA • www.dynarex.com
Made in India R220107

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DYNAREX HYDROGEL

glycerin gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-233
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	20 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POLYOXYL 60 HYDROGENATED CASTOR OIL (UNII: 02NG325BQG)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
EDETATE SODIUM (UNII: MP1J8420LU)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALLANTOIN (UNII: 344S277G0Z)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-233-02	24 in 1 CASE	05/26/2010	
1	NDC:67777-233-01	84.7 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part346	05/26/2010	

Labeler - Dynarex Corporation (008124539)

Registrant - Dynarex Corporation (008124539)

Revised: 11/2022 Dynarex Corporation