# GLYCERIN- glycerin suppository DSC Laboratories Inc.

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# **Core Values Adult Glycerin Suppositories**

## Active ingredient (in each suppository)

Glycerin 2g

# **Purpose**

Laxative

#### Uses

- for relief of occassional constipation
- this product generally produces bowel movement in 1/4 to 1 hour

### Warnings

## For rectal use only

May cause rectal discomfort or a burning sensation.

# Ask a doctor before using any laxative if you have

- abdominal pain, nausea or vomiting
- a sudden change in bowel habits lasting more than 2 weeks
- already used a laxative formore than 1 week

# Stop use and consult a doctor if you have

- rectal bleeding
- no bowel movement after using this product

These symptoms may indicate a serious condition.

# Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

# Directions-single daily dosage

adult and children 6 years and over - 1 suppository, or as directed by a doctor children 2 to under 6 years - use Child Suppositories

Insert suppository well up into rectum. Suppository need not melt completely to produce laxative action.

#### Other information

- Store container tightly closed.
- Keep away from excessive heat.

#### **Inactive ingredients**

purified water, sodium hydroxide, stearic acid

#### Questions or comments?

1-800-492-5988

#### Label

The product package shown below represents a sample of that currently in use. Additional packaging may also be available.

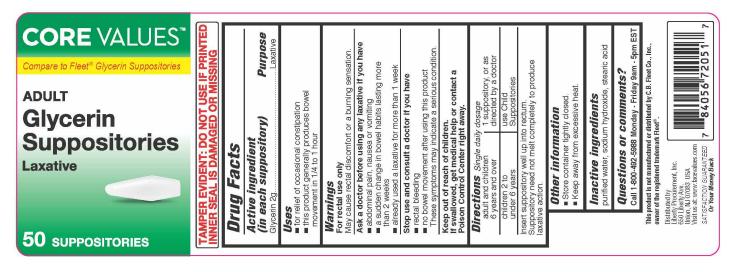
Adult Glycerin Suppositories, 50 count

Distributed by

Liberty Procurement Inc.

650 Liberty Ave.

Union, NJ 07083 USA



# glycerin suppository

**GLYCERIN** 

# Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:52316-720 Route of Administration RECTAL

#### **Active Ingredient/Active Moiety**

Ingredient Name	<b>Basis of Strength</b>	Strength
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	2 g

Inactive Ingredients			
Ingredient Name	Strength		
SODIUM HYDROXIDE (UNII: 55X04QC32I)			
STEARIC ACID (UNII: 4ELV7Z65AP)			
WATER (UNII: 059QF0KO0R)			

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:52316-720- 51	50 in 1 JAR; Type 0: Not a Combination Product	08/01/2017	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M007	08/01/2017		

# Labeler - DSC Laboratories Inc. (097807374)

# Registrant - DSC Laboratories Inc. (097807374)

Establishment					
Name	Address	ID/FEI	<b>Business Operations</b>		
DSC Laboratories Inc.		097807374	manufacture(52316-720)		

Revised: 12/2023 DSC Laboratories Inc.