GLYCERIN - glycerin suppository Chain Drug Marketing Association Inc.

Quality Choice Glycerin Supp.

Active ingredient (in each suppository)

Glycerin 2.1 g

Purpose

Laxative

Uses

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

Warnings

For rectal use only

May cause rectal discomfort of 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 for more 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

adults and children 6 years and over children 2 to under 6 years

1 suppository, or as directed by a doctor 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

GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)

Glycerin Suppositories, 50 count

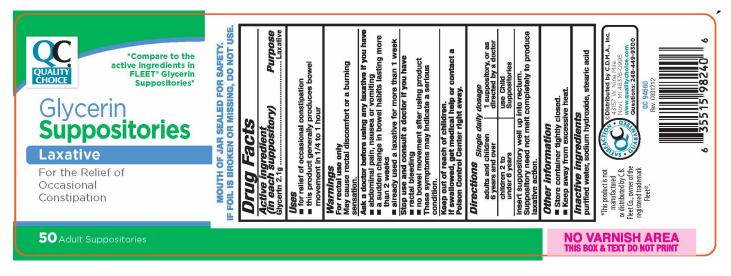
The product package shown below represents a sample of that currently in use. Additional packaging may also be available.

Glycerin Suppositories, 50 count

Distributed by Quality Choice

Novi, MI 48376-0995

www.qualitychoice.com



GLYCERIN glycerin suppository Product Information Product Type HUMAN OTC DRUG Item Code (Source) Route of Administration Active Ingredient/Active Moiety Ingredient Name Basis of Strength Strength

GLYCERIN

2.1 g

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

F	Packaging						
#	tem Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:63868-283- 50	50 in 1 JAR; Type 0: Not a Combination Product	03/15/2012				
2	NDC:63868-283- 25	25 in 1 JAR; Type 0: Not a Combination Product	03/15/2012				

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	M007	03/15/2012			

Labeler - Chain Drug Marketing Association Inc. (011920774)

Registrant - DSC Laboratories Inc. (097807374)

Establishment							
Name	Address	ID/FEI	Business Operations				
DSC Laboratories Inc.		097807374	manufacture(63868-283)				

Revised: 12/2023 Chain Drug Marketing Association Inc.