GLYCERIN - glycerin suppository
Chain Drug Marketing Association Inc.

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.

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	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

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)**	NDC:63868-283
Route of Administration	RECTAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	2.1 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

STEARIC ACID (UNII: 4ELV7Z65AP)	

Packaging

#	Item 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 not final	part334	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: 12/2019 Chain Drug Marketing Association Inc.