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	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.

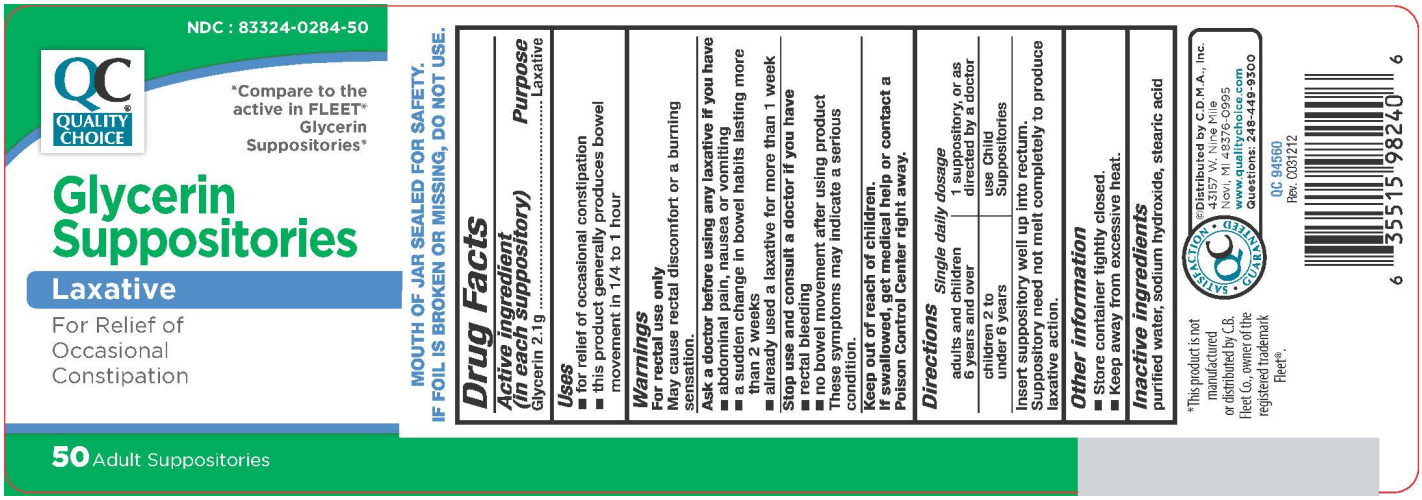
Glycerin Suppositories, 50 count

Distributed by Quality Choice

Novi, MI 48376-0995

www.qualitychoice.com

QC_50ct.jpg



GLYCERIN

glycerin suppository

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83324-284
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:83324-284-50	50 in 1 JAR; Type 0: Not a Combination Product	10/29/2025	
2	NDC:83324-284-25	25 in 1 JAR; Type 0: Not a Combination Product	10/29/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	10/29/2025	

Labeler - Chain Drug Marketing Association Inc. (011920774)

Registrant - DSC Laboratories Inc. (097807374)

Establishment

Name	Address	ID/FEI	Business Operations
DSC Laboratories Inc.		097807374	manufacture(83324-284)

Revised: 10/2025 Chain Drug Marketing Association Inc.