

**GLYCERIN - glycerin suppository**  
**Chain Drug Marketing Association Inc.**

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**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



\*Compare to the active ingredients in FLEET® Glycerin Suppositories\*

## Glycerin Suppositories

### Laxative

For the Relief of Occasional Constipation

**50** Adult Suppositories


**IF FOIL IS BROKEN OR MISSING, DO NOT USE.**

<b>Drug Facts</b>	<b>Active Ingredient (in each suppository)</b> Glycerin 2.1g ..... Laxative	<b>Purpose</b> ..... Laxative
<b>Uses</b>	■ for relief of occasional constipation ■ this product generally produces bowel movement in 1/4 to 1 hour	
<b>Warnings</b>	For rectal use only May cause rectal discomfort or a burning sensation. <b>Ask a doctor before using any laxative if you have</b> ■ 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 <b>Stop use and consult a doctor if you have</b> ■ rectal bleeding ■ no bowel movement after using product These symptoms may indicate a serious condition. <b>Keep out of reach of children.</b> <b>If swallowed, get medical help or contact a Poison Control Center right away.</b>	
<b>Directions</b>	<b>Single daily dosage</b> 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.	
<b>Other information</b>	■ Store container tightly closed. ■ Keep away from excessive heat.	
<b>Inactive Ingredients</b>	purified water, sodium hydroxide, stearic acid	

\*This product is not manufactured or distributed by C.B. Fleet Co., owner of the registered trademark Fleet®.

Manufactured by C.B. Fleet Co., Inc.  
48376 Novi, MI 48376-0995  
www.qualitychoice.com  
Questions: 248-449-9300

QC 94560  
Rev. 03/12



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**NO VARNISH AREA**  
THIS BOX & TEXT DO NOT PRINT

## 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
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)	
<b>STEARIC ACID</b> (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 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.