GLYCERIN - glycerin suppository
Chain Drug Marketing Association Inc.

Quality Choice Pediatric Glycerin Supp

Active ingredient (in each suppository)

Glycerin 1.2 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 or a burning sensation

Ask a Doctor before using any laxative if child has

- 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 child has

- rectal bleeding
- no bowel movement within 1 hour of 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

childrien 2 to under 6-1 suppository, or as directed by a doctor

children under 2 years-ask a doctor

Insert suppository well up into rectum.

Suppository need not melt completely to produce laxative action.

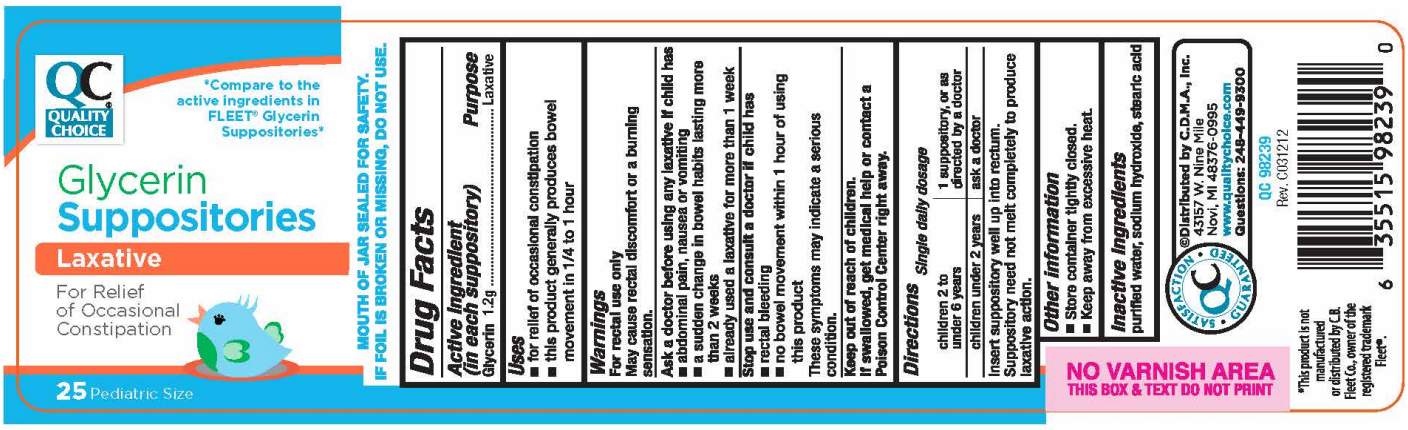
Other information

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

Inactive ingredients

purified water, sodium hydroxide, stearic acid

Pediatric Glycerin Suppositories, 25 count



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

PediatricGlycerin Suppositories, 25 count

Distributed by C.D.M.A. Inc.

43157 W. Nine Mile

Novi, MI 48376-0995

www.qualitychoice.com

QC Pediatric25ct.jpg

GLYCERIN

glycerin suppository

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-282
Route of Administration	RECTAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	1.2 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:63868-282-25	25 in 1 JAR; Type 0: Not a Combination Product	03/20/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	03/20/2012	

Labeler - Chain Drug Marketing Association Inc. (011920774)

Registrant - DSC Laboratories Inc. (097807374)

Establishment

Name	Address	ID/FEI	Business Operations
DSC Laboratories Inc.		097807374	manufacture(63868-282)

Revised: 12/2023 Chain Drug Marketing Association Inc.