GLYCERIN - glycerin suppository Chain Drug Consortium LLC

Premier Value Pediatric Glycerin Supp

Active ingredient (in each suppository)

Glycerin 1 g

Purpose

Hyperosmotic 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 laxative if child has

- abdominal pain, nausea or vomiting
- a sudden change in bowel habits lasting more than two 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

children 2 to 1 suppository or as under 6 years directed by a doctor children under 2 years ask a doctor

Insert suppository well up intorectum.

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

PremVal722.jpg

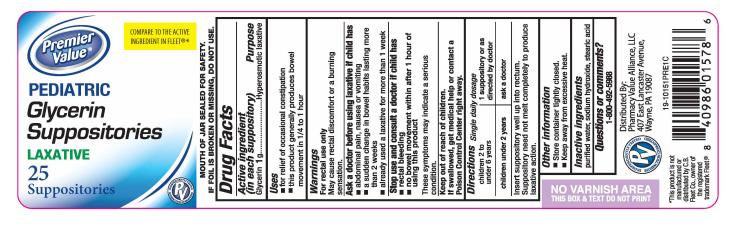
The product package shown above represents a sample of that currently in use. Additional packaging may also be available.

PediatricGlycerin Suppositories, 25 count

Distributed by Pharmacy Value Alliance, LLC

407 East Lancaster Avenue

Wayne, PA 19087



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

GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	1 g
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Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
SODIUM HYDROXIDE (UNII: 55X04QC32I)			
STEARIC ACID (UNII: 4ELV7Z65AP)			

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:68016-161- 00	25 in 1 JAR; Type 0: Not a Combination Product	03/21/2012	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	03/21/2012	

Labeler - Chain Drug Consortium LLC (101668460)

Registrant - DSC Laboratories, Inc. (097807374)

Establishment				
Name	Address	ID/FEI	Business Operations	
DSC Laboratories, Inc.		097807374	manufacture(68016-161)	

Revised: 12/2023 Chain Drug Consortium LLC