GLYCERIN- glycerin suppository
H E B

H E B Adult Glycerin Suppositories

Active ingredient (in each suppository)

Glycerin 2g

Purpose

Laxative

Uses

- for relief of occassional constipation
- this product generally produces 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 you have

- abdominal pain, nausea or vomiting
- a sudden change in bowel habits lasting more than 2 weeks
- already used a laxative formore 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 dose

adult and children 6 years and over - 1 suppository, or as directed by a doctor

chidren 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

Questions or comments?

1-800-492-5988

Label

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

Adult Glycerin Suppositories, 50 count

Distributed by H-E-B, San Antonio, TX 78204 1002

HEBAdult50.jpg

GLYCERIN

glycerin suppository

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37808-398
Route of Administration	RECTAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	2 g

Inactive Ingredients	
Ingredient Name	**Strength**
STEARIC ACID (UNII: 4ELV7Z65AP)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:37808-398-50	50 in 1 JAR; Type 0: Not a Combination Product	09/23/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	09/23/2016	

Labeler - H E B (007924756)

Registrant - DSC Laboratories Inc. (097807374)

Establishment

Name	Address	ID/FEI	Business Operations
DSC Laboratories Inc.		097807374	manufacture(37808-398)

Revised: 12/2023 H E B