STOOL SOFTENER LAXATIVE- docusate sodium capsule H E B

Drug Facts

Active ingredient (in each softgel)

Docusate Sodium 100 mg

Purpose

Stool softener laxative

Uses

- for relief of occasional constipation and irregularity
- this product generally produces bowel movement in 12 to 72 hours

Warnings

Do not use

if you are presently taking mineral oil, unless directed by a doctor

Ask a doctor before use if you have

- stomach pain
- nausea
- vomiting
- noticed a sudden change in bowel habits that last over 2 weeks

Stop use and ask a doctor if

- you have rectal bleeding or fail to have a bowel movement after use of a laxative. These could be a signs of a serious condition.
- you need to use a laxative for more than 1 week

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

• take with a glass of water

years and over	daily dose or in divided doses.
children 2 to under 12 years of age	take 1 softgel daily
children under 2 years	ask a doctor

Other information

- each softgel contains: sodium 5 mg
- store at 25°C (77°F); excursions permitted between 15-30°C (59-86°F)

Inactive ingredients

edible ink, FD&C Red #40, FD&C Yellow #6, gelatin, glycerin, polyethylene glycol, propylene glycol, purified water sorbitan, sorbitol

Questions or comments?

Call toll free: **1-877-753-3935**, Monday-Friday, 9AM-5PM EST

Principal Display Panel

Compare to Dulcolax® Stool Softener active ingredient**

Stool Softener

Laxative

Docusate Sodium 100 mg

Stool Softener Laxative

Fast, Dependable Relief of Occasional Constipation

SOFTGELS

**This product is not manufactured or distributed by Chattem, Inc., distributor of Dulcolax® Stool Softener

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

MADE WITH PRIDE AND CARE FOR H-E-B®, SAN ANTONIO, TX 78204

Product Label

Drug Facts Active ingredient Purpose (in each softgel) Docusate sodium 100 mg.. Stool softener laxative ■ for relief of occasional constipation and irregularity ■ this product generally produces bowel movement in 12 to 72 hours Warnings Do not use if you are presently taking mineral oil, unless directed by a doctor. Ask a doctor before use if you have ■ stomach pain ■ nausea ■ vomiting a sudden change in bowel habits that lasts more than 2 weeks Stop use and ask a doctor if ■ you have rectal bleeding or no bowel movement after using a laxative. These could be signs of a serious condition. ■ you need to use a laxative for more than 1 week If pregnant or breast-feeding, ask a health professional before use Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away. TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING. KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION





H-E-B Stool Softener Laxative

STOOL SOFTENER LAXATIVE

docusate sodium capsule

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:37808-862

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

Inactive Ingredients			
Ingredient Name	Strength		
FD&C RED NO. 40 (UNII: WZB9127XOA)			
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)			
GELATIN (UNII: 2G86QN327L)			
SORBITAN (UNII: 6092ICV9RU)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
WATER (UNII: 059QF0KO0R)			
SORBITOL (UNII: 506T60A25R)			
GLYCERIN (UNII: PDC6A3C0OX)			

Product Characteristics						
Color	orange	Score	no score			
Shape	OVAL	Size	12mm			
Flavor		Imprint Code	P51			
Contains						

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:37808- 862-10	1 in 1 BOX	04/30/2021				
1		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product					

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	M007	04/30/2021			
OTC Monograph Drug	M007	04/30/2021			

Labeler - H E B (007924756)

Revised: 4/2024 H E B