GENTLE STOOL SOFTENER- docusate sodium capsule, liquid filled WALGREEN COMPANY

Gentle Stool Softener

Drug Facts

Active ingredient (in each liquid-filled capsule)

Docusate Sodium 100 mg

Purpose

Stool Softner laxative

∏Uses

- relieves of occasional constipation (irregularity)
- this product generally produces a bowel movement within 12 to 72 hours

Warnings

Ask a doctor before use if you have

- stomach pain, nausea or vomittiong
- a sudden change in bowel habits that lasts more than 2 weeks

Do not useif you are presently taking mineral oil, unless told to do so by a doctor

Stop use and ask a doctor if you

- you have rectal bleeding ot no bowel movement after using this product. These could be signs of a serious condition.
- yo need to use 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.1(800)222-1222

Directions

take with a glass of water

take by mouth. Doses may be taken as a single daily dose or in divided doses.

Adults and children 12 years and overtake 1 to 3 capsules daily.

Children 2 to under 12 years of age 1 capsule daily.

Children under 2 years of age

Ask a doctor

Other information

- Store at room temperature 20-25°C (68-77°F)
- protect from excessive humidity
- do not use this product of the safety seal under the cap is torn or missing

Inactive ingredients

FD&C Blue #1*, FD&C Red #40, FD&C Yellow #6, gelatin, glycerin, polyethylene glycol, povidone, propylene glycol*, purified water, sorbitol solution *may contain this ingredient

Questions or comments?

1-800-373-6981

Distributed by: Walgreen co. 200 Wilmot rd., Deerfield, IL 60015

PRINCIPAL DISPLAY PANEL

Compare to Dulcolax ® Stool Softener active ingredient *
GENTLE STOOL SOFTENER
DOCUSATE SODIUM 100 mg / STOOL SOFTENER LAXATIVE
25 LIQUID GELS



GENTLE STOOL SOFTENER

docusate sodium capsule, liquid filled

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0363-1179

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

DOCUSATE SODIUM (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG) DOCUSATE SODIUM 100 mg

Inactive Ingredients

Ingredient Name	Strength
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POVIDONE (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SORBITOL SOLUTION (UNII: 8KW3E207O2)	

Product Characteristics					
Color	red	Score	no score		
Shape	OVAL	Size	12mm		
Flavor		Imprint Code	SD23;DX		
Contains					

P	Packaging							
#	Item Code	Package Description	Marketing Start Date	Marketing End Date				
1	NDC:0363-1179- 25	1 in 1 CARTON	03/18/2025					
1		25 in 1 BOTTLE; 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	03/18/2025		

Labeler - WALGREEN COMPANY (008965063)

Revised: 3/2025 WALGREEN COMPANY