

STOOL SOFTENER DOCUSATE SODIUM- docusate sodium capsule, gelatin coated
Magno-Humphries, Inc.

Stool Softener Docusate Sodium

Drug Facts

Active ingredient (in each softgel)

Docusate sodium 100 mg

Purpose

Stool softener

Uses

- relieves occasional constipation (irregularity)
- generally produces bowel movement in 12 to 72 hours

Warnings

Do not use

- if you are presently taking mineral oil, unless told to do so by a doctor

Ask a doctor before use if you have

- stomach pain
- nausea
- vomiting
- noticed a sudden change in bowel habits that lasts 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 signs of a serious condition.
- you need to use a stool softener 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 accidental overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

- Take only by mouth. Doses may be taken as a single daily dose, preferably in the evening, or in divided doses.

adults and children 12 years of age and older	take 1 to 3 softgels daily
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 7mg
- store at 15° to 30°C (59° to 86°F)
- protect from moisture

Inactive ingredients

FD&C red #40, FD&C yellow #6, gelatin, glycerin, polyethylene glycol, purified water, sorbitol special

Questions?

call toll-free 1-800-935-6737

Package Labeling:

NDC 54257-902-02



Stool Softener

Docusate Sodium 100 mg

Compare to the active ingredient in Colace®*
Effective, Gentle, Stimulant Free

100 SOFTGELS

DO NOT USE IF PRINTED SEAL UNDER CAP IS MISSING OR DAMAGED

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Distributed by: Magno-Humphries Labs, Tigard, OR 97223 U.S.A. L9022-100-01-3
Visit our website: www.magno-humphries.com

*This product is not manufactured or distributed by the owner of the registered trademark Colace®



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STOOL SOFTENER DOCUSATE SODIUM

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54257-902
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 YELLOW NO. 6 (UNII: H77VEI93A8)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
WATER (UNII: 059QF0K00R)	

Product Characteristics

Color	red	Score	no score
Shape	OVAL	Size	12mm
Flavor		Imprint Code	SCU1
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54257-902-02	100 in 1 BOTTLE; Type 0: Not a Combination Product	01/24/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	01/24/2020	

Labeler - Magno-Humphries, Inc. (063251433)