

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

Stool Softener Docusate Sodium

MHL NDC 54257-902-06

Stool Softener Docusate Sodium 100mg Compare to the active ingredient in Colace *

®

Effective, Gentle Stimulant Free For Comfortable Relief

1000 SOFTGELS

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

Active ingredient (in each softgel)

Docusate sodium 100mg

Purpose

Stool softener

Uses

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

Warnings

Do not use

Ask a doctor before use if you have

Stop use and ask a doctor if

ask a health professional before use. **If pregnant or breast-feeding,**

- if you are presently taking mineral oil, unless told by a doctor
- stomach pain
- nausea

- vomiting
- noticed a sudden change in bowel movements that continues over a period of 2 weeks
- you have rectal bleeding or you 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 for more than 1 week.

Keep out of reach of children.

In case of accidental overdose, get medical help or contact a Poison Control Center () right away. **1-800-222-1222**

Directions

- Take only by mouth. This dose may be taken as a single daily dose or in divided doses.
- take 1-3 softgels daily or as directed by a doctor **adults and children over 12 years of age:**
- take 1 softgel daily or as directed by a doctor **children 2 to under 12 years of age:**
- ask a doctor **children under 2 years of age:**

Other information

- sodium 7mg **each softgel contains:**
- store between 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 400, purified water, sorbitol.

Questions?

call toll-free **1-800-935-6737**

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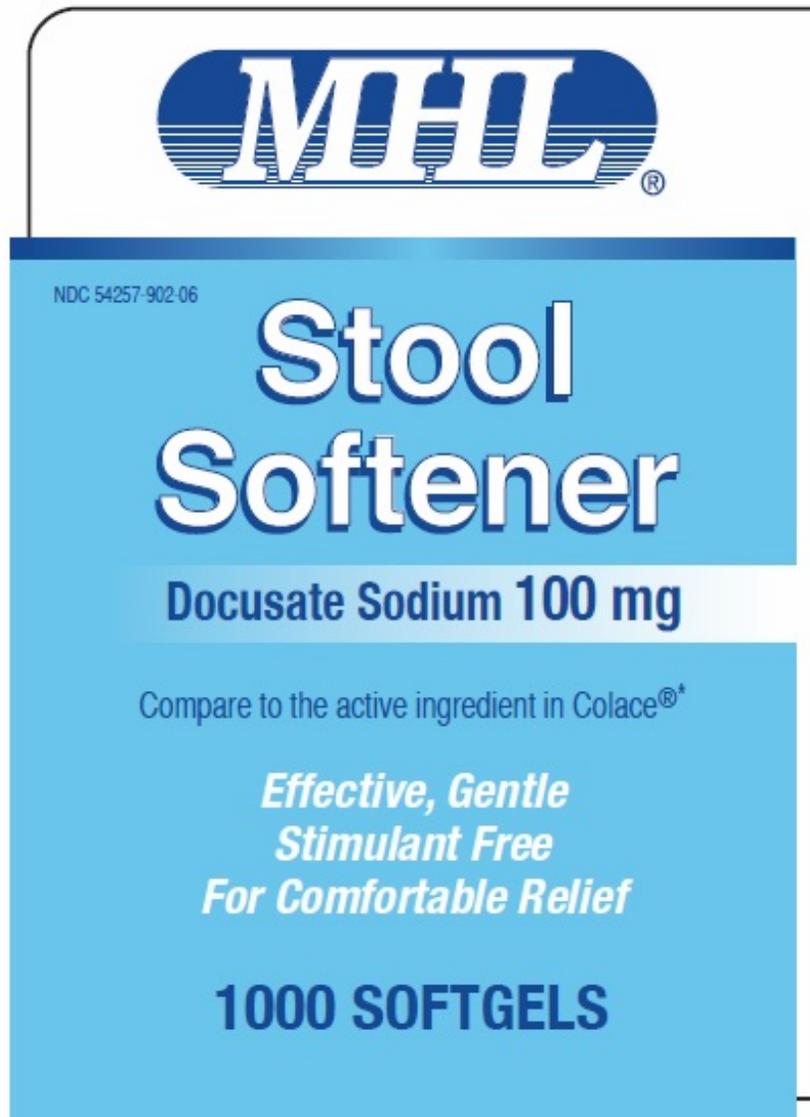
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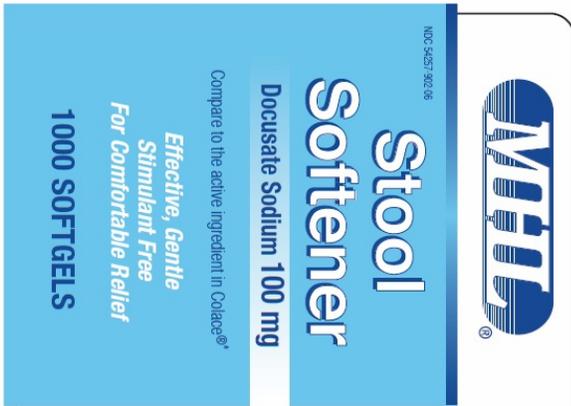
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Magno-Humphries, Inc. Visit our website: www.magno-humphries.com

Package Labeling:





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L9022-999-01-1



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docusate sodium capsule, gelatin coated

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

Ingredient Name	Strength
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	

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:54257-902-06	1000 in 1 BOTTLE; Type 0: Not a Combination Product	04/04/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	04/04/2017	

Labeler - Magno-Humphries, Inc. (063251433)

Revised: 10/2023

Magno-Humphries, Inc.