

DOCUSATE SODIUM - STOOL SOFTENER LAXATIVE- docusate sodium capsule, liquid filled

Major Pharmaceuticals

Docusate Sodium - Stool Softener Laxative

Drug Facts

Active ingredient (in each softgel)

Docusate Sodium 100 mg

Purpose

Stool softener laxative

Uses

- for relief of occasional constipation (irregularity)
- This generally produces a bowel movement within 12 to 72 hours.

Warnings

Ask a doctor before use if you have

- abdominal pain, nausea, or vomiting
- a sudden change in bowel habits that lasts over 14 days

Ask a doctor or pharmacist before use if you are presently taking mineral oil

Stop use and ask a doctor if

- you have rectal bleeding or no bowel movement after using this product.

These can 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 care professional before use.

Keep out of reach of children

Overdose Warning: In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). Prompt medical attention is critical for adults as well as children even if you do not notice any signs or symptoms.

Directions

- take softgels with a full glass (8 oz) of water

adults and
children 12

take 1 to 3 softgels daily or
as directed

CHILDREN 12 years of age and older	by a doctor. This dose may be taken as a single daily dose or in divided doses.
children 6 to under 12 years of age	take 1 softgel daily or as directed by a doctor
children under 6 years of age	ask a doctor

Other information

- **each softgel contains:** sodium 10 mg (very low sodium)
- store at 25°C (77°F) in a dry place. Avoid excessive heat 40°C (140°F).

Inactive ingredients

edible white ink, FD&C red no. 40, FD&C yellow no. 6, gelatin, glycerin, isopropyl alcohol, light mineral oil, medium chain triglycerides, polyethylene glycol, propylene glycol, purified water, sorbitol sorbitan solution

Questions or comments?

(800)616-2471

MAJOR®

Unit Dose

TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING

READ AND KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION

Distributed by:

MAJOR® PHARMACEUTICALS

Indianapolis, IN 46268

Product of UAE

Packaged and Quality Assured in the USA

Packaging



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MAJOR® NDC 0904-7183-61

Unit Dose
Docusate Sodium
Stool Softener Laxative

100 mg

Actual Size 

100 Softgels

Drug Facts (continued)

Inactive ingredients
edible white ink, FD&C red no. 40, FD&C yellow no. 6, gelatin, glycerin, isopropyl alcohol, light mineral oil, medium chain triglycerides, polyethylene glycol, propylene glycol, purified water, sorbitol sorbitan solution

Questions or comments? (866)816-2471


READ AND KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION

Distributed by:
MAJOR PHARMACEUTICALS
Indianapolis, IN 46268
Product of UAE
Packaged and Quality Assured in the USA
CT0904718361
REV.01-03/2022

MAJOR® NDC 0904-7183-61

Unit Dose
Docusate Sodium
Stool Softener Laxative

100 mg

Actual Size 

100 Softgels

Rev. 07/21 M-173 Re-order No. 701293

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Lot:
Exp:

DOCUSATE SODIUM - STOOL SOFTENER LAXATIVE			
docusate sodium capsule, liquid filled			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0904-7183
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 RED NO. 40 (UNII: WZB9127XOA)		
	FD&C YELLOW NO. 6 (UNII: H77VEI93A8)		
	GELATIN, UNSPECIFIED (UNII: 2G86QN327L)		
	GLYCERIN (UNII: PDC6A3C0OX)		

ISOPROPYL ALCOHOL (UNII: ND2M416302)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
SORBITAN (UNII: 6O92ICV9RU)	

Product Characteristics

Color	red	Score	no score
Shape	OVAL	Size	12mm
Flavor		Imprint Code	777
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0904-7183-61	10 in 1 CARTON	09/04/2021	
1		10 in 1 BLISTER PACK; 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	09/04/2021	

Labeler - Major Pharmaceuticals (191427277)

Revised: 12/2024

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