# SENNA S- sennosides and docusate sodium tablet, coated Medline Industries, LP

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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## 989 Senna S Sennosides 8.6mg and Docusate Sodium 50mg

## Active ingredient (in each tablet)

- Docusate sodium 50 mg
- Sennosides 8.6 mg

### **Purpose**

Stool softener laxative

Stimulant laxative

#### Uses

- relieves occasional constipation (irregularity)
- generally produces a bowel movement in 6 -12 hours

## Warnings

#### Do not use

- this product if you are presently taking mineral oil, unless directed by a doctor
- laxative products for longer than 1 week unless directed by a doctor

# Ask a doctor before use if you have

- stomach pain
- nausea
- vomiting
- a sudden change in bowel movements that persists 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.

# 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 preferably at bedtime or as directed by a doctor
- if you do not have a comfortable bowel movement by the second day, increase dose by one tablet (do not exceed maximum dosage) or decrease dose until you are comfortable

	_	maximum dosage
adults and children 12	2 tablets	4 tablets
years of age and over	once a day	twice a day
children 6 to	1 tablet	2 tablets
under 12 years	once a day	twice a day
children 2 to	1/2 tablet	1 tablet
under 6 years	once a day	twice a day
children under 2 years	ask a doctor	ask a doctor

#### Other information

- each tablet contains: calcium 20 mg, sodium 6 mg (VERY LOW SODIUM)
- store at 25°C (77°F); excursions permitted between 15-30°C (59-86°F)

### Inactive ingredients

carnauba wax, colloidal silicon dioxide, croscarmellose sodium, dibasic calcium phosphate dihydrate, D&C yellow#10 aluminum lake, FD&C yellow#6 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, sodium benzoate, stearic acid, titanium dioxide

#### **Ouestions or comments?**

Call toll free **1-800-MEDLINE (633-5463)** Monday-Friday 9AM-5PM CST

# **Manufacturing Information**

Manufactured for:
Medline Industries, LP
Three Lakes Drive, Northfield, IL 60093 USA
Made in USA with domestic and foreign materials
www.medline.com
1-800-MEDLINE (633-5463)

REF: OTCM00011 V1 RE22HND

# Package Label



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Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Centerright away. hypromellose, magnesium stearate, microcrystalline 4 tablets 2 tablets 1 tablet silicon dioxide, croscarmellose sodium, dibasic calcium phosphate dihydrate, D&C yellow#10 cellubse, polyethylene glycol, sodium benzoate, aluminum lake, FD& C yello w#6 aluminum lake, if pregnant or breast-feeding, ask a health Hinge Area Questions or comments?
Call toll free 1-800-MEDLINE (633-5463)
Montaw-Friday 9AM-5PM (ST once a day once a day once a day 1/2 tablet starting 2 tablets dosage 1 tablet Drug Facts (continued) years of age and over children 6 to under 12 years children 2 to adults and children 12 children under 2 years Other information you are comfortable. 15-30°C (59-86°F1 under 6 years П Directions 

## **SENNAS**

sennosides and docusate sodium tablet, coated

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53329-989	
Route of Administration	ORAL			

Active Ingredient/Active Moiety				
Ingredient Name	<b>Basis of Strength</b>	Strength		
DOCUSATE SODIUM (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG)	DOCUSATE SODIUM	50 mg		
SENNOSIDES (UNII: 3FYP5M0IJX) (SENNOSIDES - UNII:3FYP5M0IJX)	SENNOSIDES	8.6 mg		

Inactive Ingredients			
Ingredient Name	Strength		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
CARNAUBA WAX (UNII: R12CBM0EIZ)			
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			
CALCIUM PHOSPHATE, DIBASIC, DIHYDRATE (UNII: O7TSZ97GEP)			
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)			
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)			
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)			

HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics				
Color	orange	Score	score with uneven pieces	
Shape	ROUND	Size	10mm	
Flavor		Imprint Code	TCL;081	
Contains				

Packaging						
	# Item (	Code	Package Description		Marketing Start Date	Marketing End Date
	1 NDC:533 989-62		1 BOTTLE, PLASTIC; Type 0: Not a ination Product		05/22/2018	



# **Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	05/22/2018	

# Labeler - Medline Industries, LP (025460908)

# Registrant - Medline Industries, LP (025460908)

Revised: 12/2022 Medline Industries, LP