SENNA S- docusate sodium and sennosides tablet Marlex Pharmaceuticals Inc

Senna S

SENNOSIDES AND DOCUSATE SODIUM-sennosides, docusate sodium tabletMarlexPharmaceuticals, Inc.

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.

SennaS

Drug Facts

Active ingredient (in each tablet)

Docusate Sodium 50mg Sennosides 8.6 mg

Purpose

Stool softener Stimulant laxative

Uses

- relieves occasional constipation (irregularity)
- generally causes 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 doctorif

you have rectal bleeding or fail to have a bowel movement after use if 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 does by one tablet (do not exceed maximum dosage) or decrease does until you are comfortable

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

Other information

- each tablet contains:calcium 4mg, sodium 8mg
- store at 20°C to 25°C (68°F to 77°F) excursions permitted between 15°30°C (59°-86°F)

Inactive ingredients

Colloidal silicon dioxide, croscarmellose sodium, D&C Yellow #10 Aluminum Lake, FD&C Yellow #6 Aluminum Lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, sodium benzoate, titanium dioxide

*Compare to the active ingredients in SENOKOT-S®

PRINCIPAL DISPLAY PANEL

NDC 10135-0669-01 Senna S 100 TABLET



ACTUAL SIZE • DIE LINE DOES NOT PRINT

Drug Facts (continued)	Ask a doctor before use if you have stomach pain In rausea In vomiting In a sudden change in bowel movements that persists over 2 weeks Stop use and ask a doctor if you have rectal bleeding or fall 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 Po ison Contro (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.	starting maximum dosage dosage	hildren 12 2 tablets 4 tablets and over once a day twice a day	1 tablet 2 tablets ars once a day twice a day	1/2 tablet 1 tablet once a day twice a day	er 2 years ask a doctor ask a doctor	Other information ■ each tablet contains: calcium 4 mg, sodium 8 mg ■ store at 20°C to 25°C (68°F to 77°F); excursions permitted between 15-30°C (59-86°F)	Inactive ingredients colloidal silion dioxide, α oscarmellose sodium, D&C Yellow #10 Aluminum Lake, FD&C Yellow #6 Aluminum Lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, sodium berzoate, titanium dioxide
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SENNA S

docusate sodium and sennosides tablet

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10135-669		
Route of Administration	ORAL				

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

Inactive Ingredients	
Ingredient Name	Strength

SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics					
Color	orange	Score	no score		
Shape	ROUND	Size	9mm		
Flavor		Imprint Code	S6		
Contains					

P	Packaging							
#	Item Code	Package Description	Marketing Start Date	Marketing End Date				
1	NDC:10135-669- 01	1 in 1 CARTON	10/01/2018					
1		100 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	10/01/2018				

Labeler - Marlex Pharmaceuticals Inc (782540215)

Revised: 11/2023 Marlex Pharmaceuticals Inc