SENNA-S- docusate sodium and sennosides tablet Spirit Pharmaceuticals LLC

Docusate Sodium 50mg, Sennosides 8.6mg Tablets Drug Facts

Active ingredients (in each tablet)

Docusate sodium 50 mg Sennosides 8.6 mg

Purpose

Stool Softner

Laxative

Uses

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

Warnings

Do not use

- If you are now 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
- noticed a sudden change in bowel habits that lasts over a period of 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 may indicate 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. 1(800)222-1222

Directions

• take preferably at bedtime or as directed by a doctor

children 6 to under 12 years
children 2 to under 6 years
children under 2 years

1 tablet once a day 1/2 tablet once a day 1 tablet twice a day ask a doctor

2 tablets twice a day ask a doctor

Other information

- each tablet contains: calcium 19.92 mg, sodium 5.61 mg
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)

Inactive ingredients

Colloidal silicon dioxide, croscarmellose sodium, dicalcium phosphate, D&C yellow No. 10, FD&C Yellow No. 6, hypromellose, magnesium stearate, microcrystalline cellulose, maltodextrin, polyethylene glycol-400, purified water, sodium benzoate, stearic acid, titanium dioxide

Questions or comments?

1-888-333-9792

PDP

Cabinet:

Stool Softener & Laxative DOCUSATE SODIUM 50 mg SENNOSIDES 8.6 mg **75 TABLETS**

Compare to the active ingredients in Senokot-S®*

Stool Softener & Laxative

Docusate Sodium 50mg Sennosides 8.6mg

75 Tablets

SENNA-S

docusate sodium and sennosides tablet

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:68210-4103

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	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
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)	
ANHYDROUS DIBASIC CALCIUM PHOSPHATE (UNII: L11K75P92J)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	

MALTODEXTRIN (UNII: 7CVR7L4A2D)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	orange	Score	no score
Shape	ROUND (biconvex)	Size	10mm
Flavor		Imprint Code	S35
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68210- 4103-1	75 in 1 BOTTLE; Type 0: Not a Combination Product	07/02/2020	
2	NDC:68210- 4103-2	200 in 1 BOTTLE; Type 0: Not a Combination Product	07/02/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	07/02/2020	

Labeler - Spirit Pharmaceuticals LLC (179621011)

Revised: 12/2023 Spirit Pharmaceuticals LLC