

SENNA-S LAXATIVE- docusate sodium, sennosides tablet
New World Imports, 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.

Active ingredients (in each tablet)

Docusate Sodium 50 mg

Sennosides 8.6 mg

Purpose

Stool softener

Laxative

Uses

- relieves occasional constipation (irregularity).
- generally causes bowel movement within 6 to 12 hours.

Warnings

Do not use:

☐ ☐☐ This product is 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.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

- taken preferably at bedtime or as directed by a doctor

Age	Starting Dose	Maximum Dosage
adults and children 12 years of age and older	2 tablets once a day	4 tablets twice a day
children 6 to under 12 years of age	1 tablet once a day	2 tablets twice a day
children 2 to 6 years of age	1/2 tablet once a day	1 tablet twice a day

children under 2 years

ask a doctor

ask a doctor

Other information

- each tablet contains: **sodium 6 mg/tablet** LOW SODIUM
- each tablet contains: **calcium 20 mg/tablet**
- store at 15°-30°C (59°-86°F), in well closed containers
- do not use if imprinted safety seal under cap is broken or missing
- *This product is not manufactured or distributed by Purdue Pharma L.P., owner of the registered trademark Senokot S.

Inactive ingredients

carnauba wax*, croscarmellose sodium, DandC Yellow 10 Aluminum Lake, dibasic calcium phosphate dihydrate, FDandC Blue 2 Aluminum Lake*, FDandC Red 40 Aluminum Lake*, FDandC Yellow 6 Aluminum Lake, hypromellose*, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol*, purified water*, sodium benzoate*, stearic acid, titanium dioxide

*may also contain

NDC 51824-026-01

Compare to the active ingredients in SENOKOT-S[®]*

Senna-S Laxative

Natural Vegetable Laxative

▶ Stool Softener Plus Laxative

For Gentle, Overnight Relief of Constipation

60

Docusate sodium, 50 mg

Sennosides, 8.6 mg Tablets

DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

Drug Facts

Active ingredients (in each tablet) Purpose
 Docusate Sodium 50 mg.....Stool softener
 Sennosides 8.6 mg.....Stimulant laxative

Uses
 ■ relieves occasional constipation (irregularity)
 ■ generally produces 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 care professional before use.
Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Drug Facts (continued under label)

0 85317 00561 0

Distributed by:
 NWI, Inc., 160 Athens Way
 Nashville, TN 37228
 LB1164
 R0912

Lot No.:
 Exp. Date:

PEEL HERE

Drug Facts (continued)

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

Age	Starting dosage	Maximum dosage
Adults and children 12 years and over	2 tablets once a day	4 tablets twice a day
Children 6 to under 12 years	1 tablet once a day	2 tablets twice a day
Children 2 to under 6 years	1/2 tablet once a day	1 tablet twice a day
Children under 2 years	ask a doctor	ask a doctor

Other information
 ■ each tablet contains: calcium 20 mg, sodium 6 mg (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.

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SENNAS-S LAXATIVE

docusate sodium, sennosides tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51824-026
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
CARNAUBA WAX (UNII: R12CBM0EIZ)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OPIR32D61U)	
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL (UNII: 532B59J990)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	orange	Score	no score
Shape	ROUND	Size	9mm
Flavor		Imprint Code	TCL081
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51824-026-01	60 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/01/2012	08/01/2020

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	12/01/2012	08/01/2020

