

SENNA S- docusate sodium sennosides tablet
QUALITY CHOICE (Chain Drug Marketing Association)

Drug Facts

Active ingredients (in each tablet)

Docusate sodium 50 mg
Sennosides 8.6 mg

Purpose

Stool softener
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 continues 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 (1-800-222-1222) right away.

Directions

- take preferably at bedtime or as directed by a doctor

age	starting dosage	maximum dosage
adults and children 12 years of age or 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 30 mg
- **each tablet contains:** sodium 6 mg LOW SODIUM
- store at 25°C (77°F); excursions permitted between 15-30°C (59-86°F)

Inactive ingredients

croscarmellose sodium, dibasic calcium phosphate dihydrate, FD&C yellow #6 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, silicon dioxide, talc, titanium dioxide

Questions or comments?

Call **1-800-935-2362** Monday-Friday 9AM-5PM EST

Principal Display Panel

**Compare to the Active Ingredients in Senokot-S®

Stool Softener Plus Laxative

Stool Softener • Laxative

Docusate Sodium 50 mg | Sennosides 8.6 mg

Provides Gentle Relief of: Occasional Constipation

Tablets

**This product is not manufactured or distributed by Atlantis Consumer Healthcare Inc., distributor Senokot-S®.

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION

Distributed by CDMA, Inc.

Novi, MI 48375

Product Labeling

NDC 83324-135-60

****Compare to the
Active Ingredients
in Senokot-S®**



Stool Softener

Plus Laxative

Stool Softener • Laxative

Docusate Sodium 50 mg | Sennosides 8.6 mg
Provides Gentle Relief of: Occasional Constipation

60 Tablets

Actual Size



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TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

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Drug Facts (continued)

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Docusate sodium 50 mg

Stool softener
Laxative



Distributed by CDMA, Inc.
Novi, MI 48375
www.qualitychoice.com
Questions: 800-935-2362



PLD-F599A
FC009138

Lot No.:

Exp. Date:

QUALITY CHOICE Stool Softener plus Laxative

SENNA S

docusate sodium sennosides tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83324-135
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
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
ALUMINUM OXIDE (UNII: LMI26O6933)	

Product Characteristics

Color	orange	Score	no score
Shape	ROUND	Size	10mm
Flavor		Imprint Code	PSD22
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-135-60	1 in 1 BOX	07/31/2024	
1		60 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
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OTC Monograph Drug	M007	07/31/2024	
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Labeler - QUALITY CHOICE (Chain Drug Marketing Association) (011920774)

Revised: 5/2024

QUALITY CHOICE (Chain Drug Marketing Association)