SENOKOT-S- standardized senna concentrate and docusate sodium tablet
Atlantis Consumer Healthcare, Inc.

Senokot-S
(standardizedsenna concentrate and docusate sodium)

Drug Facts

Active ingredients (in each tablet)

Docusate sodium 50mg

Sennosides 8.6mg

Purpose

Stool softener

Laxative

Uses

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

Warnings

Do not use

- if you are now taking mineral oil, unless directed by adoctor
- laxative products for longer than 1 week unless directedby a doctor

Ask a doctor before use if you have

- stomach pain
- nausea
- vomiting
- noticed a sudden change in bowel habits that continues overa period of 2 weeks

Stop using and ask a doctor if youhave rectal bleeding or fail to have a bowel movement after use ofa laxative. These may indicate a serious condition.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reachof children.

In case ofoverdose, get medical help or contact a Poison Control Center rightaway.

Directions

- take preferably at bedtime or as directed by a doctor

age	startingdosage	maximumdosage
adults and children 12 years of age and over	2 tablets oncea day	4 tablets twicea 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

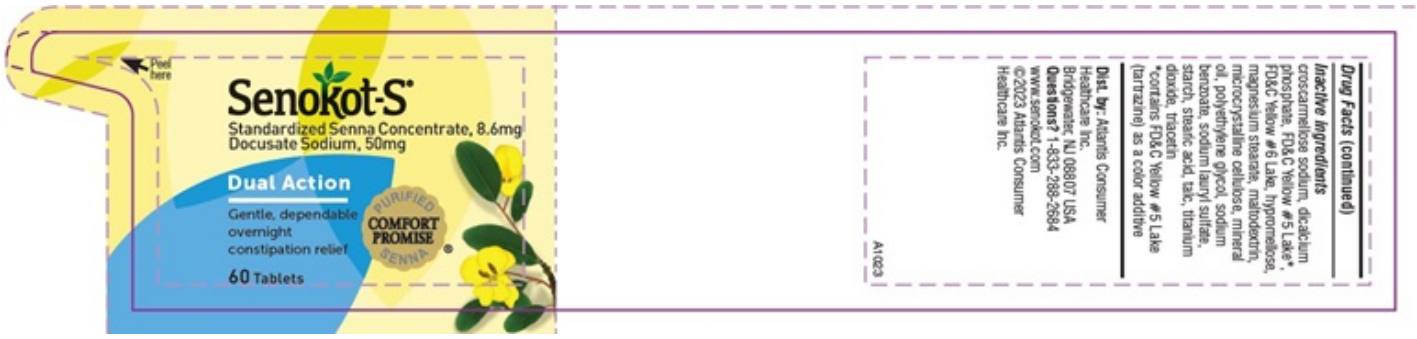
- each tablet contains: **calcium 7 mg, sodium 4 mg VERYLOW SODIUM**
- store at 25°C (77°F); excursions permitted between 15°-30°C(59°-86°F)

Inactive ingredients

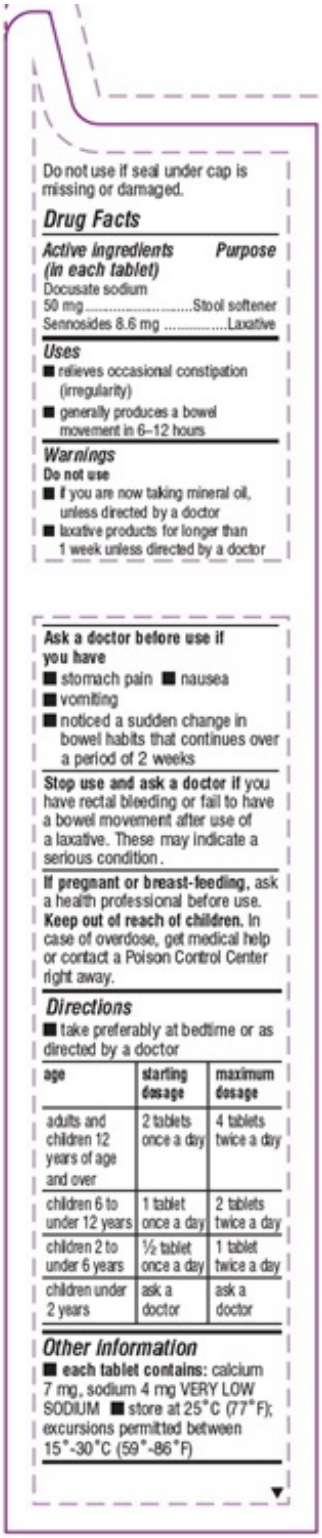
Inactive ingredients croscarmellose sodium, dicalcium phosphate, FD&C Yellow#5 Lake*, FD&C Yellow #6 Lake, hypromellose, magnesium stearate,maltodextrin, microcrystalline cellulose, mineral oil, polyethyleneglycol, sodium benzoate, sodium lauryl sulfate, starch, stearic acid,talc, titanium dioxide, triacetin

A1023

Senokot-S 60 Tablets Label
NDC: 67618-310-60



Senokot-S 60 Tablets Leaflet
NDC: 67618-310-60



Do not use if seal under cap is missing or damaged.

Drug Facts

Active ingredients (in each tablet)	Purpose
Docusate sodium 50 mg	Stool softener
Sennosides 8.6 mg	Laxative

Uses

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- 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 right away.

Directions

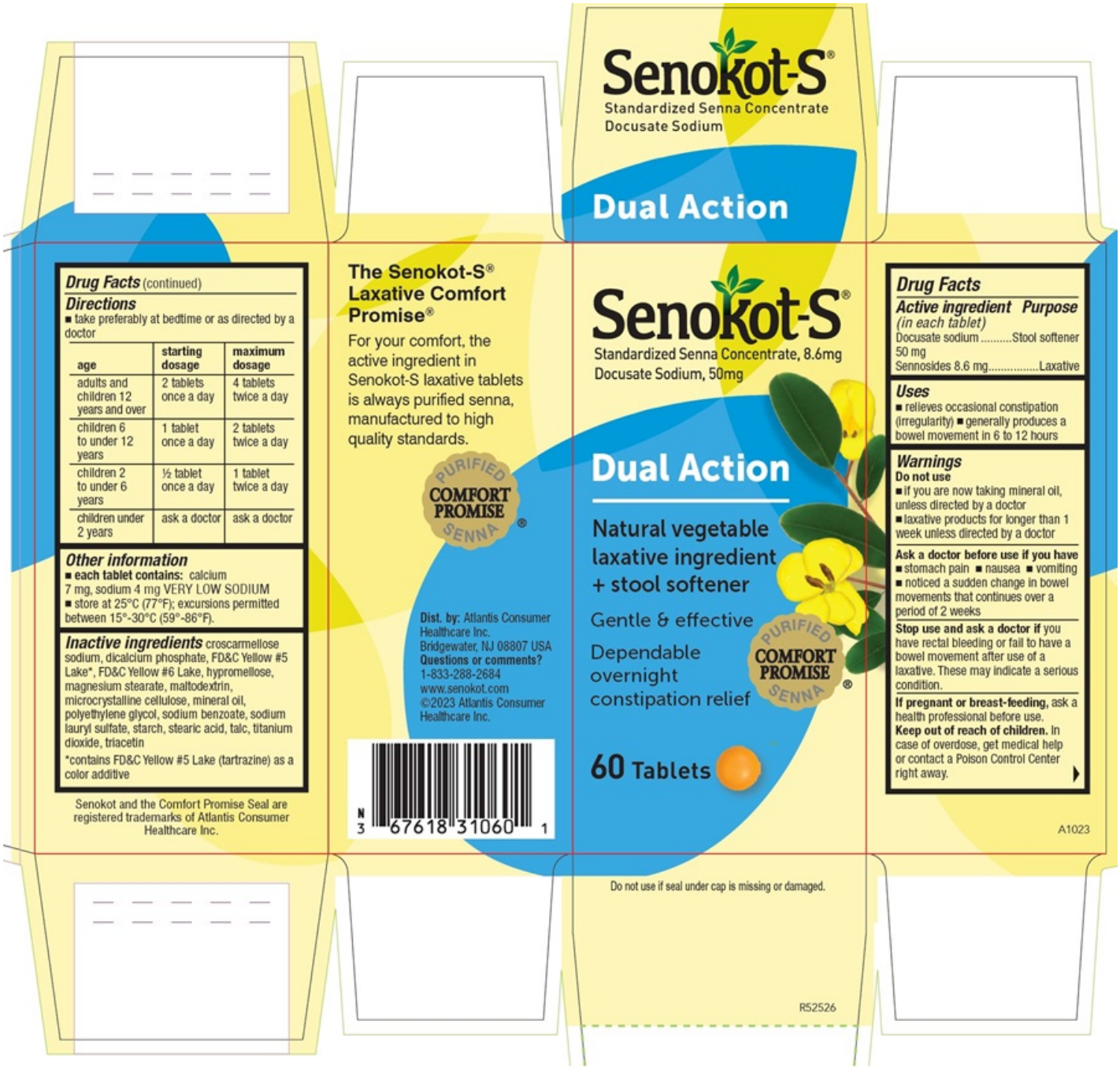
- take preferably at bedtime or as directed by a doctor

age	starting dosage	maximum dosage
adults and children 12 years of age 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	½ tablet once a day	1 tablet twice a day
children under 2 years	ask a doctor	ask a doctor

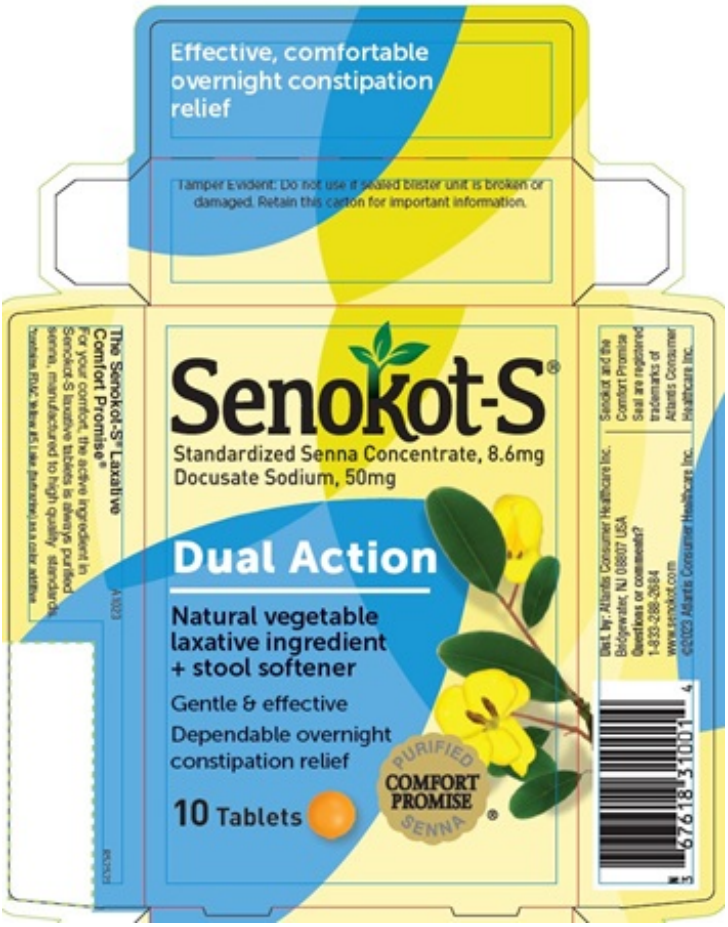
Other information

- **each tablet contains:** calcium 7 mg, sodium 4 mg VERY LOW SODIUM
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)

Senokot-S 60 Tablets Carton
NDC: 67618-310-60



Senokot-S 10 Tablets Carton
NDC: 67618-310-01


Effective, comfortable overnight constipation relief
Tamper Evident: Do not use if sealed blister unit is broken or damaged. Retain this carton for important information.
The Senokot-S® Laxative Comfort Promise®
For your comfort, the active ingredient in Senokot-S laxative tablets is always purified senna, manufactured to high quality standards.
Senokot-S®
Standardized Senna Concentrate, 8.6mg
Docusate Sodium, 50mg
Dual Action
Natural vegetable laxative ingredient + stool softener
Gentle & effective
Dependable overnight constipation relief
10 Tablets
PURIFIED COMFORT PROMISE SENNA
Senokot and the Comfort Promise Seal are registered trademarks of Atlantis Consumer Healthcare Inc.
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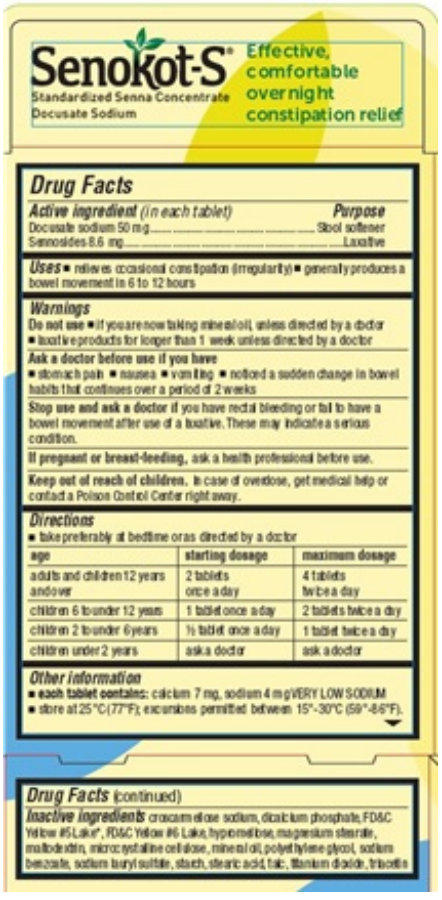
Dist. by: Atlantis Consumer Healthcare Inc. Bridgewater, NJ 08807 USA
Questions or comments?
1-833-288-3684
www.senokot.com
3 67618 31001 4


Senokot-S®
Standardized Senna Concentrate
Docusate Sodium
Effective, comfortable overnight constipation relief
Drug Facts
Active ingredient (in each tablet) Purpose
Docusate sodium 50 mg Stool softener
Sennosides 8.6 mg Laxative
Uses ■ relieves occasional constipation (irregularity) ■ generally produces a bowel movement in 6 to 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 right away.
Directions
■ take preferably at bedtime or as directed by a doctor
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 | ½ 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 7 mg, sodium 4 mg VERY LOW SODIUM
■ store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F).
Drug Facts (continued)
Inactive ingredients croscarmellose sodium, dicalcium phosphate, FD&C Yellow #5 Lake*, FD&C Yellow #6 Lake, hypromellose, magnesium stearate, maltodextrin, microcrystalline cellulose, mineral oil, polyethylene glycol, sodium benzoate, sodium lauryl sulfate, starch, stearic acid, talc, titanium dioxide, triacetin

SENOKOT-S

standardized senna concentrate and docusate sodium tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67618-310
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**
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
CALCIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: L11K75P92J)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
MINERAL OIL (UNII: T5L8T28FGP)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	

Product Characteristics			
Color	ORANGE	**Score**	no score
Shape	ROUND	**Size**	10mm
Flavor		**Imprint Code**	P
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:67618-310-01	1 in 1 CARTON	10/01/1974	
1		10 in 1 BLISTER PACK; Type 0: Not a Combination		

1		Product		
2	NDC:67618-310-30	1 in 1 CARTON	10/01/1974	
2		30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:67618-310-60	1 in 1 CARTON	10/01/1974	
3		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
OTC Monograph Drug	M007	10/01/1974	

Labeler - Atlantis Consumer Healthcare, Inc. (118983925)

Registrant - Atlantis Consumer Healthcare, Inc. (118983925)

Revised: 12/2023 Atlantis Consumer Healthcare, Inc.