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

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***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

- 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 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 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?
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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: 12/2025 Atlantis Consumer Healthcare, Inc.