**SENNA S- docusate sodium and sennosides tablet**
**Marlex Pharmaceuticals Inc**

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**Senna S**

**SENNOSIDES AND DOCUSATE SODIUM-sennosides, docusate sodium tabletMarlexPharmaceuticals, 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.*

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**SennaS**

***Drug Facts***

***Active ingredient (in each tablet)***

Docusate Sodium 50mg Sennosides 8.6 mg

***Purpose***

Stool softener Stimulant laxative

***Uses***

- relieves occasional constipation (irregularity)
- generally causes 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 doctorif**

you have rectal bleeding or fail to have a bowel movement after use if a laxative. These could be signs of 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
- if you do not have a comfortable bowel movement by the second day, increase does by one tablet (do not exceed maximum dosage) or decrease does until you are comfortable

| **age** | **starting dosage** | **maximum dosage** |
|---|---|---|
| adults and children 12 years of age and older | 2 tablets once a day | 4 tablets twice a day |
| children 6 to 12 years | 1 tablet once a day | 2 tablets twice a day |
| children 2 to 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 4mg, sodium 8mg
- store at 20°C to 25°C (68°F to 77°F) excursions permitted between 15°30°C (59°-86°F)

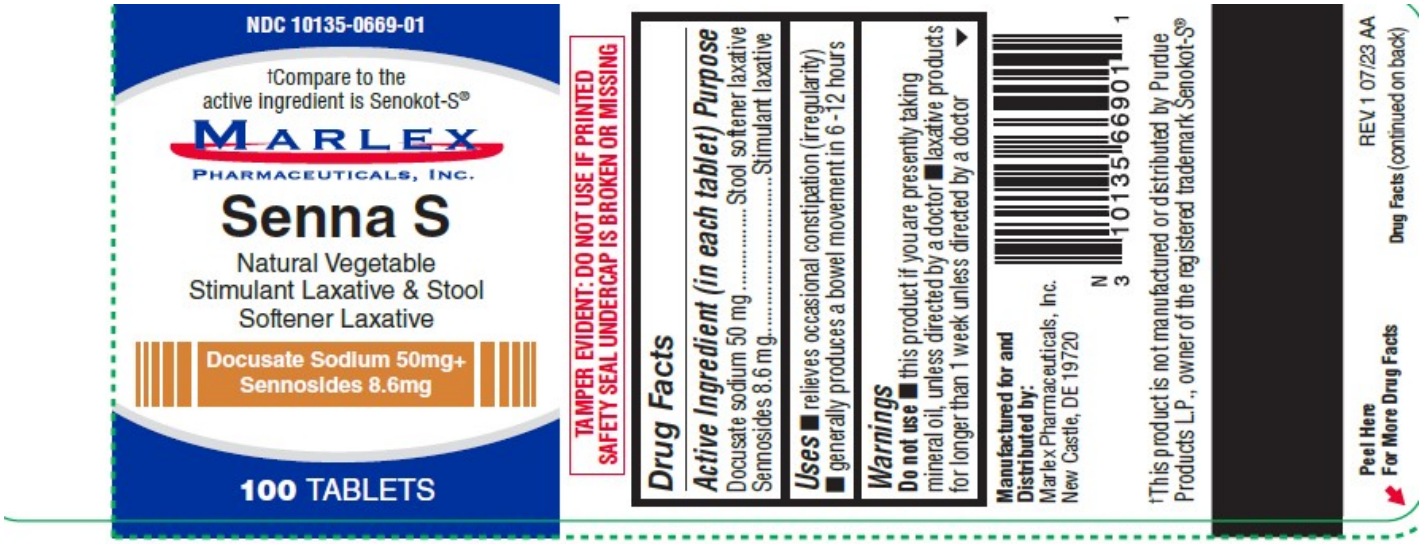
***Inactive ingredients***

Colloidal silicon dioxide, croscarmellose sodium, D&C Yellow #10 Aluminum Lake, FD&C Yellow #6 Aluminum Lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, sodium benzoate, titanium dioxide

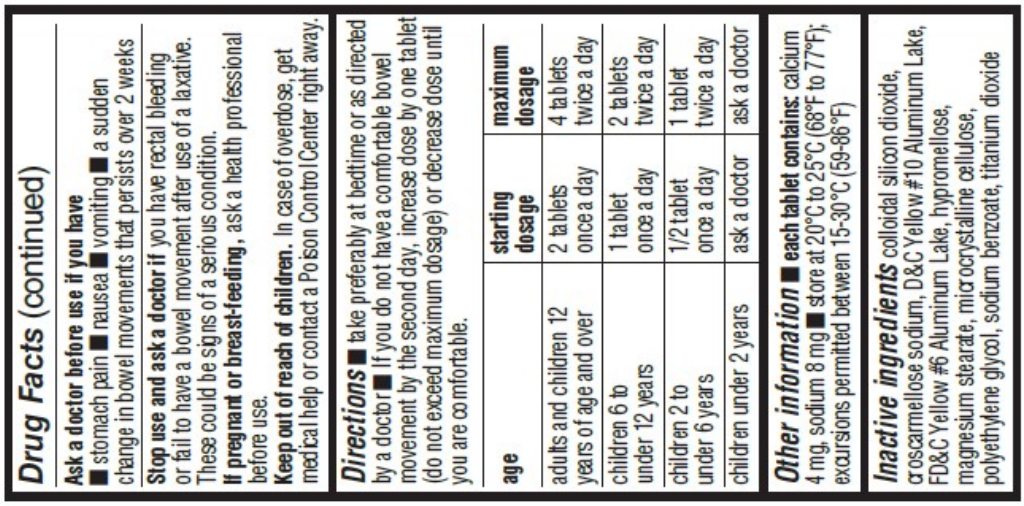
*Compare to the active ingredients in SENOKOT-S®

**PRINCIPAL DISPLAY PANEL**

NDC 10135-0669-01
Senna S
100 TABLET




ACTUAL SIZE • DIE LINE DOES NOT PRINT




# SENNA S

docusate sodium and sennosides tablet

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:10135-669 |
| **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** |

| | |
|---|---|
| **SILICON DIOXIDE** (UNII: ETJ7Z6XBU4) | |
| **CROSCARMELLOSE SODIUM** (UNII: M28OL1HH48) | |
| **D&C YELLOW NO. 10** (UNII: 35SW5USQ3G) | |
| **FD&C YELLOW NO. 6** (UNII: H77VEI93A8) | |
| **HYPROMELLOSES** (UNII: 3NXW29V3WO) | |
| **MAGNESIUM STEARATE** (UNII: 70097M6I30) | |
| **CELLULOSE, MICROCRYSTALLINE** (UNII: OP1R32D61U) | |
| **POLYETHYLENE GLYCOL, UNSPECIFIED** (UNII: 3WQ0SDW1A) | |
| **SODIUM BENZOATE** (UNII: OJ245FE5EU) | |
| **TITANIUM DIOXIDE** (UNII: 15FIX9V2JP) | |

## Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | orange | **Score** | no score |
| **Shape** | ROUND | **Size** | 9mm |
| **Flavor** | | **Imprint Code** | S6 |
| **Contains** | | | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:10135-669-01 | 1 in 1 CARTON | 10/01/2018 | |
| **1** | | 100 in 1 BOTTLE; 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 | 10/01/2018 | |

## Labeler - Marlex Pharmaceuticals Inc (782540215)

Revised: 10/2024 Marlex Pharmaceuticals Inc