

**SENNA-S- sennosides 8.6mg and docusate sodium 50mg tablet, film coated
Pharbest Pharmaceuticals, Inc.**

Drug Facts

Active Ingredients (in each tablet)

Sennosides from Senna Concentrate 8.6mg

Docusate Sodium 50mg

Purpose

Laxative

Stool Softner

Uses

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

Warnings

Do not use

- laxative products for longer than 1 week unless directed by a doctor
- if you are taking mineral oil, 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 the use of a laxative. These may indicate a serious condition.

If pregnant or breast feeding,

ask a healthcare professional before use.

Keep out of reach of children.

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

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	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 **10 mg of calcium, sodium 5 mg**
- store at 25°(77°F); excursions permitted between 15°-30°C (59°-86°F)
- **Tamper Evident:** Do not use if safetyseal under cap is broken or missing

Inactive ingredients

crosscarmellose Sodium, D&C Yellow# 10, dicalcium phosphate, FD&C Yellow #6, magnesium stearate, microcrystalline cellulose, polyvinyl alcohol, polyethylene glycol, sodium benzoate, talc, titanium dioxide

Questions or comments?

(866) 562-2756 Mon-Fri 8 AM to 4 PM EST

PHARBEST

NDC 16103-378-08

Manufactured in the USA

****Compare to the active ingredients in Senokot-S®***

SENNAS

Sennosides 8.6mg &

Docosate Sodium 50mg

Natural Vegetable Laxative

Ingredient Plus Stool Softner

100 TABLETS

Compare to the active ingredients in Senokot-S

SENNA-S

Sennosides 8.6mg & Docusate Sodium 50mg

Natural Vegetable Laxative Ingredient Plus Stool Softener

100 TABLETS

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If pregnant or breast feeding , ask a healthcare professional before use. Keep out of reach of children. In case of overdose, get medical help or	

Peel here for more drug facts



Manufactured by: Pharmed Pharmaceuticals, Inc., Farmingdale, NY 11735

Drug Facts (contd.) contact a Poison Control Center immediately.	Directions ■ take preferably at bedtime or as directed by a doctor	starting dosage	maximum dosage
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Original 02/19
The product is not manufactured or distributed by Purdue Pharma L.P., owner of the registered trademark, Senokot-S.
STOP PEELING

SENNA-S

sennosides 8.6mg and docusate sodium 50mg tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:16103-378
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)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
CALCIUM PHOSPHATE, DIBASIC, DIHYDRATE (UNII: O7TSZ97GEP)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	

POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	orange (ORANGE COLOR)	Score	no score
Shape	ROUND (ROUND TABLET)	Size	10mm
Flavor		Imprint Code	PH32
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:16103-378-07	60 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/05/2018	
2	NDC:16103-378-11	1000 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/05/2018	
3	NDC:16103-378-08	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/21/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	11/05/2018	

Labeler - Pharbest Pharmaceuticals, Inc. (557054835)

Registrant - Pharbest Pharmaceuticals, Inc. (557054835)

Establishment

Name	Address	ID/FEI	Business Operations
Pharbest Pharmaceuticals, Inc.		557054835	manufacture(16103-378) , analysis(16103-378) , pack(16103-378) , label(16103-378)

Revised: 12/2025

Pharbest Pharmaceuticals, Inc.