

**SENNA-S- sennosides 8.6mg and docusate sodium 50mg tablet, film coated
Proficient Rx LP**

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.

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)

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

NDC 71205-970-30

Manufactured in the USA

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

SENNA-S

Sennosides 8.6mg &

Docosate Sodium 50mg

Natural Vegetable Laxative

Ingredient Plus Stool Softner

30 TABLETS

Repackaged & Relabeled by:

Proficient Rx LP

Thousand Oaks, CA 91320



NDC 71205-970-30



Made in the USA

Lot #:00000
Exp. 00/00/00
SN# MASTER

Senna & Docusate 8.6mg/50mg

#30 Tablets

Each tablet contains: Sennosides from Senna Concentrate 8.6mg Laxative/ Docusate Sodium 50mg Stool Softner

Orange, round, unscored tablet with imprint code PH32

Product ID: QS097030

Mfr. By: Pharbest Pharmaceuticals, Inc., Farmingdale NY 11735

Store at 25°C (77°F)

Keep medication out of the reach of children

Packaged By: Proficient Rx LP
Thousand Oaks, CA 91320



SENNAS

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

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71205-970(NDC:16103-378)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SENNOSIDES (UNII: 3FYP5M0IIX) (SENNOSIDES - UNII:3FYP5M0IIX)	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)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (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:71205-970-30	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2020	
2	NDC:71205-970-60	60 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2020	
3	NDC:71205-970-90	90 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2020	
4	NDC:71205-970-00	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2020	
5	NDC:71205-970-55	500 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2020	
6	NDC:71205-970-11	1000 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	11/05/2018	

Labeler - Proficient Rx LP (079196022)

Establishment

Name	Address	ID/FEI	Business Operations
Proficient Rx LP		079196022	REPACK(71205-970) , RELABEL(71205-970)

Revised: 6/2020

Proficient Rx LP