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

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

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starting dosage maximum dosage
age
Adults and
            2 tablets once a 4 tablets twice a
children 12
            day
                            day
vears and
over
            1 tablet once
Children 6
                            2 tablets twice
to under 12 a day
                            a day
vears
Children 2
            1/2 tablet once 1 tablet twice
to under 6 a day
                            a day
vears
Children
            ask a doctor ask a doctor
under 2
years
```

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 &

Docusate Sodium 50mg

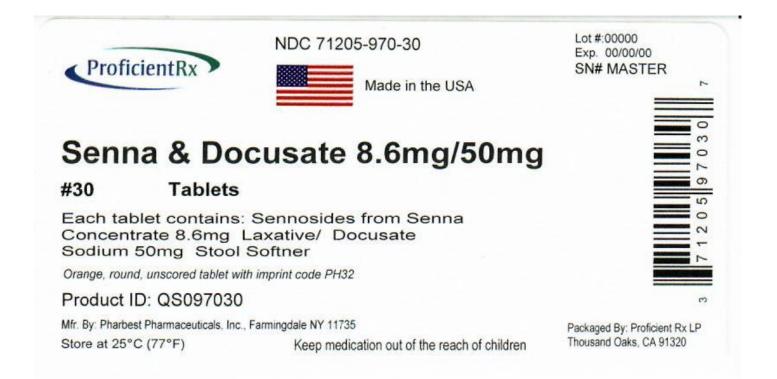
Natural Vegetable Laxative

Ingredient Plus Stool Softner

30 TABLETS

Repackaged & Relabele by:

Proficient Rx LP Thousand Oaks, CA 91320



SENNA-S

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: 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: M280L1HH48)				
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

#	ltem 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	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
OTC Monograph Drug	M007	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: 10/2024

Proficient Rx LP