# 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.

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# **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	2 tablets once a	4 tablets twice a day		
children 12	day			
years and over				
Children 6	1 tablet once a day	2 tablets twice a day		
to under 12				
years				
Children 2	1/2 tablet once	1 tablet twice a day		
to under 6 years a day				
Children under 2	ask a doctor			
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** 

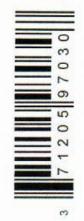
Thousand Oaks, CA 91320



NDC 71205-970-30



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

#### **SENNA-S**

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

#### **Product Information**

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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	
SENNO SIDES (UNII: 3FYP5M0 IJX) (SENNO SIDES - UNII: 3FYP5M0 IJX)	SENNOSIDES	8.6 mg	
DOCUSATE SODIUM (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG)	DOCUSATE SODIUM	50 mg	

Inactive Ingredients				
Ingredient Name	Strength			
CROSCARMELLOSE SODIUM (UNII: M28 OL1HH48)				
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)				
DIBASIC CALCIUM PHO SPHATE DIHYDRATE (UNII: O7TSZ97GEP)				
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
MICRO CRYSTALLINE CELLULO SE (UNII: OP1R32D61U)				
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)				
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)				
SO DIUM BENZO ATE (UNII: OJ245FE5EU)				
TALC (UNII: 7SEV7J4R1U)				
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)				

Product Characteristics			
Color	orange (ORANGE COLOR)	Score	no score
Shape	ROUND (ROUND TABLET)	Size	10 mm
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