

**SENNA-S- docusate sodium and sennosides tablet, film coated**  
**Contract Pharmacy Services-PA**

*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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**SENNA-S Tablets**

**DRUG FACTS**

**ACTIVE INGREDIENT (IN EACH TABLET)**

Docusate sodium 50 mg

Sennosides 8.6 mg

**PURPOSE**

Stool softener & Stimulant laxative

**USES**

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

**WARNINGS**

**Do not use**

- this product if you are presently taking mineral oil, unless directed by a doctor
- laxative products for more 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 habits that persists over 2 weeks

**Stop use and ask a doctor if you have**

- have rectal bleeding
- fail to have a bowel movement after the use of 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 dose by one tablet (do not exceed maximum dosage); or decrease dose until you are comfortable
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age	starting dose	maximum dose
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: calcium 20 mg, sodium 6 mg (LOW SODIUM)
- TAMPER-EVIDENT: Do not use this product if foil seal on back of blister is cut, torn, broken or missing
- store at 25°C (77°F) excursions permitted between 15°-30°C (59°-86°F)

#### **INACTIVE INGREDIENTS**

carnauba wax, colloidal silicon dioxide, croscarmellose sodium, dibasic calcium phosphate dihydrate, D&C yellow #10 aluminum lake, FD&C yellow #6 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, sodium benzoate, stearic acid, tapioca starch1, tartaric acid1, titanium dioxide

1may contain these ingredients

#### **QUESTIONS OR COMMENTS?**

call 1-888-838-2872, weekdays, 8 AM-5 PM Eastern Time

Distributed by:

GOLDLINE LABORATORIES, INC.

Sellersville, PA 18960 Dist. 1999 0110REV 89

192B R1009

Repackaged by:

Contract Pharmacy Services-PA


125 Titus Ave Suite 200

Warrington, PA 18976 USA

Original--08/2010--NJW

#### **Package Label**

**SENNAS-**  
8.6/50MG TAB #30



48991002770920

LOT# SAMPLE  
ID: OTC 817  
EXP: 03/12

GOLDLINE

USUAL DOSAGE:  
SEE INSERT


STORAGE  
CONDITIONS: SEE  
INSERT  
(STORE AT  
CONTROLLED ROOM  
TEMPERATURE)

FILLED BY: \_\_\_\_\_

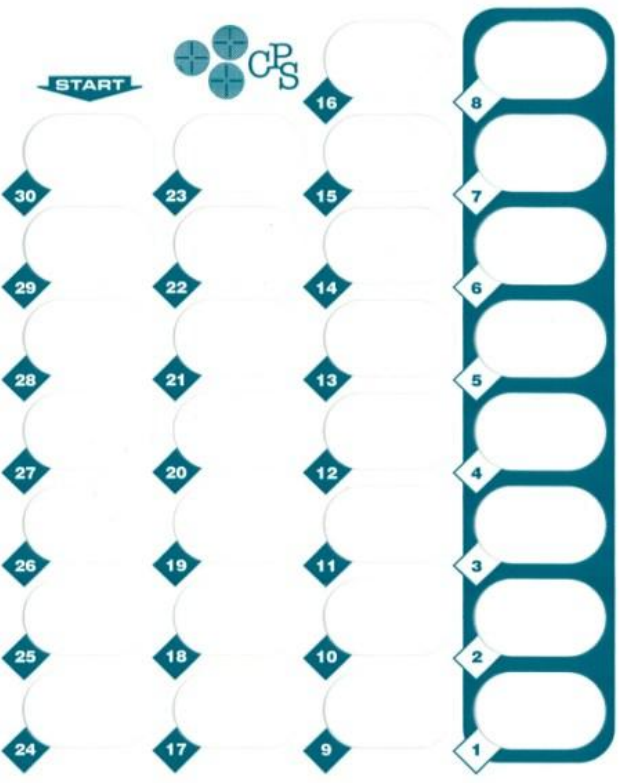
CHECKED BY: \_\_\_\_\_


Med. \_\_\_\_\_  
Strength \_\_\_\_\_  
Exp. Date \_\_\_\_\_  
Mfg. \_\_\_\_\_  
Lot No. \_\_\_\_\_

Packaged By: Contract Pharmacy Services-PA  
125 Titus Avenue, Suite #200, Warrington, PA 18976



Contract Pharmacy Services





CAUTION: This package NOT CHILD RESISTANT. Store this and all medications out of reach of children.

## SENNAS-S

docusate sodium and sennosides tablet, film coated

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:67046-670(NDC:0182-1113)
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>DOCUSATE SODIUM</b> (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG)	DOCUSATE SODIUM	50 mg
<b>SENNOSIDES</b> (UNII: 3FYP5M0IJX) (SENNOSIDES - UNII:3FYP5M0IJX)	SENNOSIDES	8.6 mg

### Inactive Ingredients

Ingredient Name	Strength
CARNAUBA WAX (UNII: R12CBM0EIZ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	

DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)
HYPROMELLOSE (UNII: 3NXW29V3WO)
MAGNESIUM STEARATE (UNII: 70097M6I30)
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)
SODIUM BENZOATE (UNII: OJ245FE5EU)
STEARIC ACID (UNII: 4ELV7Z65AP)
STARCH, TAPIOCA (UNII: 24SC3U704I)
TARTARIC ACID (UNII: W4888I119H)
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)
ALUMINUM OXIDE (UNII: LM26O6933)

Product Characteristics			
Color	YELLOW	Score	no score
Shape	ROUND	Size	8mm
Flavor		Imprint Code	8
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67046-670-30	30 in 1 BLISTER PACK		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	08/17/2010	

**Labeler** - Contract Pharmacy Services-PA (945429777)