SENNA/DOCUSATE SODIUM - docusate sodium and sennosides tablet, film coated
Physicians Total Care, Inc.

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.

Senna-S

TAMPER-EVIDENT: Do not use this product if imprinted foil seal over the mouth of the bottle is cut, torn, broken or missing.

Drug Facts

Active ingredients (in each tablet)	***Purposes***
Docusate sodium 50 mg	Stool softener
Sennosides 8.6 mg	Stimulant laxative

	Purposes
Docusate sodium 50 mg	Stool softener
Sennosides 8.6 mg	Stimulant laxative

Uses

- relieves occasional constipation (irregularity)
- generally causes 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 longer 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 rectal bleeding
- fail to have a bowel movement after 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 overodse, 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

age	starting dosage	maximum dosage
adults and children 12 years and older	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)
- store at room temperature 15° to 30°C (59° to 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 starch[1], tartaric acid[1], titanium dioxide

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1 may contain these ingredients

Questions?

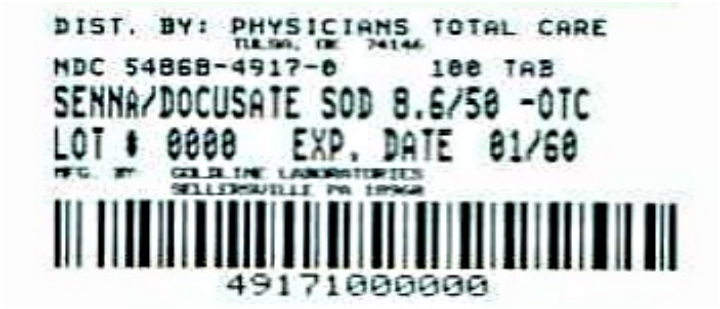
call **1-888-838-2872,** weekdays, 8 AM-5 PM Eastern Time

†This product is not manufactured or distributed by Purdue Products L.P., owner of the registered trademark Senokot-S®

Distributed by: **GOLDLINE LABORATORIES, INC.**
Sellersville, PA 18960 Dist. 1999 0110REV 89

Additional barcode labeling by:
Physicians Total Care, Inc.
Tulsa, Oklahoma 74146

PRINCIPAL DISPLAY PANEL - 100 Tablet Bottle



NDC 54868-4917-0

TAMPER-EVIDENT

Senna/Docusate Sodium Tablets
(brand of standardized senna concentrate equivalent to 8.6 mg sennosides and docusate sodium 50 mg)

Natural Vegetable Stimulant
Laxative And Stool Softener

100 TABLETS

Compare to active ingredients of Senokot-S® Tablets†

SENNA/DOCUSATE SODIUM

docusate sodium and sennosides tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54868-4917(NDC:0182-1113)
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Docusate sodium (UNII: F05Q2T2JA0) (Docusate - UNII:M7P27195AG)	Docusate sodium	50 mg
Sennosides (UNII: 3FYP5M0IJX) (Sennosides - UNII:3FYP5M0IJX)	Sennosides	8.6 mg

Inactive Ingredients	
Ingredient Name	**Strength**
carnauba wax (UNII: R12CBM0EIZ)	
silicon dioxide (UNII: ETJ7Z6XBU4)	
croscarmellose sodium (UNII: M28OL1HH48)	
dibasic calcium phosphate dihydrate (UNII: O7TSZ97GEP)	
D&C yellow NO. 10 (UNII: 35SW5USQ3G)	
FD&C yellow NO. 6 (UNII: H77VEI93A8)	
hypromelloses (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: LMI26O6933)	

Product Characteristics

Color	ORANGE	**Score**	no score
Shape	ROUND (bi-convex)	**Size**	10mm
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54868-4917-0	100 in 1 BOTTLE, PLASTIC		
2	NDC:54868-4917-1	60 in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part334	09/11/2003	

Labeler - Physicians Total Care, Inc. (194123980)

Establishment

Name	Address	ID/FEI	Business Operations
Physicians Total Care, Inc.		194123980	relabel

Revised: 4/2012 Physicians Total Care, Inc.