SENNA-S- senna and docusate sodium tablets, 8.6 mg and 50 mg tablet, coated

HealthLife of USA LLC

Senna-S - Senna and Docusate Sodium Tablets, 8.6 mg & 50 mg

Drug Facts

Active ingredients Purpose (in each tablet) Purpose

Docusate sodium 50 mg.....Stool softner

Sennosides 8.6 mg.....Laxative

Uses

- relieves occasional constipation (irregularity)
- generally produces bowel movement in 6-12 hours

Warnings

Do not use

- if you are now 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
- noticed a sudden change in bowel habits that lasts over 2 weeks

Stop use and ask a doctor if

you have rectal bleeding or fail to have a bowel movement after use of a laxative. These may indicate 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

- adults and children 12 years of age or older- starting dosage: 2 tablets once a day, maximum dosage: 4 tablets twice a day
- children 6 to under 12 years- starting dosage:1 tablet once a day, maximum dosage:2 tablets twice a day
- children 2 to under 6 years- starting dosage:1/2 tablet once a day, maximum dosage:1 tablet twice a day

children under 2 years- starting dosage:ask a doctor, maximum dosage:ask a doctor

Other information

- each tablet contains: calcium 21 mg
- each tablet contains: sodium 3 mg VERY LOW SODIUM
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- read all product information before using
- TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

Inactive ingredients

croscarmellose sodium, colloidal silicon dioxide, dicalcium phosphate, D&C Yellow # 10, FD&C Yellow # 6, hypromellose, microcrystalline cellulose, magnesium stearate, maltodextrin, polyethylene glycol, stearic acid, titanium dioxide

Questions or comments?

Call toll free 1-844-832-1138 Monday through Friday 9AM – 5PM EST or www.healthlifeofusa.com

PRINCIPAL DISPLAY PANEL

Compare to the Active Ingredients inSenokot-S®

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

Senna and Docusate Sodium Tablets, 8.6 mg and 50 mg



SENNA-S

senna and docusate sodium tablets, 8.6 mg and 50 mg tablet, coated

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69517-131
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)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CALCIUM PHOSPHATE, DIBASIC, DIHYDRATE (UNII: O7TSZ97GEP)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
POLYETHYLENE GLYCOL 1000 (UNII: U076Q6Q621)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	orange	Score	no score
Shape	ROUND	Size	10mm
Flavor		Imprint Code	S35
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69517-131- 25	25 in 1 BOX	04/08/2016	
1		2 in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:69517-131- 50	50 in 1 BOX	04/08/2016	
2		2 in 1 POUCH; Type 0: Not a Combination Product		
3	NDC:69517-131- 02	2 in 1 POUCH	04/08/2016	
3		2 in 1 POUCH; Type 0: Not a Combination Product		
4	NDC:69517-131- 10	1000 in 1 BOTTLE	04/08/2016	
4	NDC:69517-131- 05	500 in 1 BOTTLE		
4	NDC:69517-131- 24	24 in 1 BOTTLE		
4		1 in 1 CARTON; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	03/07/2016	

Labeler - HealthLife of USA LLC (079656178)

Establishment				
Name	Address	ID/FEI	Business Operations	
Elysium Pharmaceutical Ltd.		915664486	manufacture(69517-131)	

Revised: 10/2023 HealthLife of USA LLC