SENEXON-S- docusate sodium -sennosides tablet, coated NCS HealthCare of KY, Inc dba Vangard Labs

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.

Senexon S 50mg 8 6mg

Active ingredients (in each tablet)

Docusate sodium 50 mg Sennosides 8.6 mg

Purposes

Stool Softener

Stimulant Laxative

Uses

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

Warnings

Do not use

- if you are taking mineral oil
- for longer than one week
- when abdominal pain, nausea or vomiting are present

Ask a doctor before use if you

have a sudden change in bowel habits that lasts over two weeks

Ask a doctor or pharmacist before use if you are

taking any other drug. Take this product two or more hours before or after other drugs. Laxatives may affect how other drugs work.

Stop use and ask a doctor if

- you fail to have a bowel movement after use of this product
- you have rectal bleeding

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
- if you do not have a comfortable bowel movement by the second day, increase dose by one tablet (do not exceed maxiumum dosage)

age	starting dosage	maximum dosage
adults and children 12 years and older	2 tablets once daily	4 tablets twice daily
children 6 to under 12 years	1 tablet once daily	2 tablets twice daily
children 2 to under 6 years	1/2 tablet once daily	1 tablet twice daily
children under 2 years	ask a doctor	ask a doctor

Other information

- each tablet contains: calcium 20mg, sodium 6 mg (LOW SODIUM)
- store at 20° 25°C (68° 77°F)

*Rugby Laboratories is not affiliated with the owner of the registered trademark Senokot-S[®]. 192R R-14 Rev. 02/13

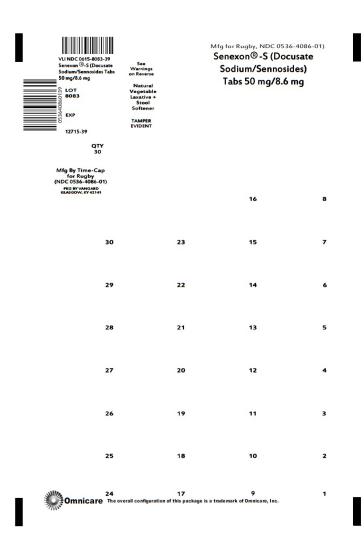
Questions or comments?

1-800-645-2158

Inactive ingredients

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

Principal Display Panel



Received:		(SEE USP CONTROLL	D ROOM TEMPERATURE)
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STORE AT 20°- 25 °C (68°- 77°F)
(SEE USP CONTROLLED ROOM TEMPERATURE)

SENEXON-S

docusate sodium -sennosides tablet, coated

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0615-8083(NDC:0536-4086)
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 A AND B (UNII: 1B5FPI42EN) (SENNOSIDES A AND B - UNII:1B5FPI42EN)	SENNOSIDES A AND B	8.6 mg		

Inactive Ingredients		
Ingredient Name	Strength	
CARNAUBA WAX (UNII: R12CBM0EIZ)		
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)		
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)	
HYPROMELLOSES (UNII: 3NXW29 V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICRO CRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)	

Product Characteristics				
Color	ORANGE	Score	no score	
Shape	ROUND	Size	9 mm	
Flavor		Imprint Code	TCL081	
Contains				

ı	P	ackaging			
ı	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:0615-8083- 39	30 in 1 BLISTER PACK; Type 0: Not a Combination Product	12/20/2016	08/31/2020

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part334	03/16/2011	08/31/2020

Labeler - NCS HealthCare of KY, Inc dba Vangard Labs (050052943)

Establishment				
Name	Address	ID/FEI	Business Operations	
NCS HealthCare of KY, Inc dba Vangard Labs		050052943	REPACK(0615-8083)	

Revised: 1/2017 NCS HealthCare of KY, Inc dba Vangard Labs