

# **SENNOSIDES AND DOCUSATE SODIUM- sennosides and docusate sodium tablet**

**American Health Packaging**

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**Sennosides and Docusate Sodium Tablets**  
**Natural Vegetable Laxative with Stool Softener**  
**0462201/0621**

## **Active ingredient (in each tablet)**

Docusate Sodium	50 mg
Sennosides	8.6 mg

## **Purpose**

Stool softener  
Laxative

## **Uses**

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

## **Warnings**

**Do not use** for more than one week unless directed by a doctor.

## **Ask a doctor before use if you**

- have abdominal pain, nausea or vomiting
- are taking mineral oil
- have noticed a sudden change in bowel habits that lasts over two 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**

- do not exceed 8 tablets in 24 hours
- Adults and children 12 years of age and older take 2 tablets once a day preferably at

bedtime; increase as needed, or as directed by a doctor; the maximum dose should be 4 tablets in the morning and 4 tablets at bedtime

- Children under 12 years ask a doctor

### **Other information**

- each tablet contains: **calcium 20 mg, sodium 3 mg**
- store at controlled room temperature
- **Keep this and all drugs out of reach of children.**
- **FOR YOUR PROTECTION:** Do not use if blister is torn or broken.

### **Inactive ingredients**

cellulose, croscarmellose sodium, dicalcium phosphate, FD&C yellow #6 lake, hypromellose, magnesium stearate, PEG, silica, talc, titanium dioxide.

### **PACKAGING INFORMATION**

The drug product contained in this package is from NDC # 57896-555, Geri-Care Pharmaceuticals Corp.

Distributed by:

American Health Packaging, Columbus, Ohio 43217

762201

0462201/0621A

**Package/Label Display Panel - Carton - 8.6 mg/50 mg**

NDC 60687-622-01

**Sennosides\* and Docusate Sodium Tablets**  
Natural Vegetable Laxative with Stool Softener

\*Standardized Senna Concentrate

**8.6 mg/50 mg**

100 Tablets (10 x 10)



(01) 0 03 60687 622 01 9

**Drug Facts**

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Docusate Sodium 50 mg.....	Stool softener
Sennosides 8.6 mg.....	Laxative

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**Drug Facts (continued)**

**Warnings (continued)**

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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.

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### Package/Label Display Panel - Blister - 8.6 mg/50 mg



Sennosides and Docusate  
Sodium Tablet  
Laxative/Stool Softener

**8.6 mg/50 mg**

**SENNOSIDES AND DOCUSATE SODIUM**  
sennosides and docusate sodium tablet

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:60687-622(NDC:57896-555)
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

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

### Inactive Ingredients

Ingredient Name	Strength
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>CROSCARMELLOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>ANHYDROUS DIBASIC CALCIUM PHOSPHATE</b> (UNII: L11K75P92J)	
<b>FD&amp;C YELLOW NO. 6</b> (UNII: H77VEI93A8)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	

### Product Characteristics

<b>Color</b>	orange	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	9mm
<b>Flavor</b>		<b>Imprint Code</b>	PSD22
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60687-622-01	100 in 1 BOX, UNIT-DOSE	08/03/2021	
1	NDC:60687-622-11	1 in 1 BLISTER PACK; Type 0: Not a Combination Product		



### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	08/03/2021	

**Labeler** - American Health Packaging (929561009)

### Establishment

Name	Address	ID/FEI	Business Operations
American Health Packaging		929561009	repack(60687-622)

Revised: 11/2024

American Health Packaging