

**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**  
**0462201/0825**

***Active ingredients(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**

- 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 or vomiting
- 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. Very low sodium.
- store at 20°-25°C (68°-77°F); excursions permitted between 15°-30°C (59°-86°F).
- **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.

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

Distributed by:

American Health Packaging, Columbus, Ohio 43217

Product of India

762201

0462201/0825

**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

**8.6 mg/50 mg**

100 Tablets (10 x 10)



**Drug Facts**

**Active ingredients (in each tablet).....Purpose**  
Docusate Sodium 50 mg.....Stool softener  
Sennosides 8.6 mg.....Laxative

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• this product generally produces a bowel movement in 6 to 12 hours

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Natural Vegetable Laxative with Stool Softener

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**Drug Facts (continued)**

**Warnings (continued)**

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NDC 60687- 622-01

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

**8.6 mg/50 mg**

**100 Tablets (10 x 10)**

**Drug Facts**

**Active ingredients (in each tablet).....Purpose**

Docusate Sodium 50 mg.....Stool softener

Sennosides 8.6 mg.....

# Laxative

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

Product of India

Distributed by:  
American Health Packaging, Columbus, Ohio 43217

762201  
0462201/0825

## 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

**Product Type**

HUMAN OTC DRUG

**Item Code (Source)**

NDC:60687-622(NDC:57896-555)

Route of Administration ORAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>SENNOSIDES</b> (UNII: 3FYP5M0JJX) (SENNOSIDES - UNII:3FYP5M0JJX)	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: 8/2025

American Health Packaging