SENNA-PLUS- sennosides and docusate sodium tablet Geri-Care Pharmaceutical Corp

hst 455b (555)

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

- 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
- 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

Age	Starting Dose	Maximum Dose
adults and children	2 tablets once a day preferably at	4 tablets in the
12 years of age and	bedtime; increase as needed, or as	morning and 4 tablets

older	directed by a doctor	at bedtime
children under 12	ask a doctor	
years	ask a doctor	

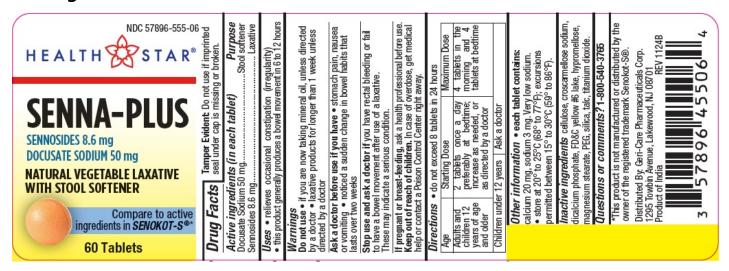
Other information

- each tablet contains: calcium 20 mg, sodium 3 mg
- store at 20° to 25°C (68° to 77°F); excursions permitted between 15° to 30°C (59° to 86°F).

Inactive ingredients

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

Package Label



SENNA-PLUS

sennosides and docusate sodium tablet

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:57896-555	
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

CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
CALCIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: L11K75P92J)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TALC (UNII: 7SEV7J4R1U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

Product Characteristics					
Color	orange	Score	no score		
Shape	ROUND	Size	9mm		
Flavor		Imprint Code	PSD22		
Contains					

P	Packaging Packag				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:57896-555- 01	100 in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2019		
2	NDC:57896-555- 06	60 in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2019		
3	NDC:57896-555- 10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2019		
4	NDC:57896-555- 20	200 in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2019		

Marketing Information				
Marketing Application Number or Monograph Marketing Start Marketing Category Citation Date Date				
OTC Monograph Drug	505G(a)(3)	05/01/2019		

Labeler - Geri-Care Pharmaceutical Corp (611196254)

Registrant - Geri-Care Pharmaceutical Corp (611196254)

Revised: 11/2024 Geri-Care Pharmaceutical Corp