SENNA-PLUS- sennosides and docusate sodium tablet
Preferred Pharmaceuticals Inc.

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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 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

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

	directed by a doctor
children under 12 years	ask a doctor

Other information

- each tablet contains: **calcium 20 mg, sodium 3 mg**
- store at controlled room temperature
  Tamper Evident: Do not use if imprinted seal under cap is missing or broken.
- product of India

Repackaged By: Preferred Pharmaceuticals Inc.

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:68788-8780(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
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
ANHYDROUS DIBASIC CALCIUM PHOSPHATE (UNII: L11K75P92J)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
HYPROMELLOSE, UNSPECIFIED (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			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-8780-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	12/11/2024	
2	NDC:68788-8780-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	12/11/2024	
3	NDC:68788-8780-8	120 in 1 BOTTLE; Type 0: Not a Combination Product	12/11/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	12/11/2024	

Labeler - Preferred Pharmaceuticals Inc. (791119022)

Registrant - Preferred Pharmaceuticals Inc. (791119022)

Establishment

Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals Inc.		791119022	REPACK(68788-8780)