SENNA SYRUP- sennosides syrup Pioneer Life Sciences, LLC

Senna Syrup

Active Ingredient (per 5 mL teaspoon)

Sennosides 8.8 mg

Purpose

Laxative

Uses

- Relieves occasional constipation (irregularity)
- Generally causes bowel movement in 6 to 12 hours

Warnings

Do not use laxative products for longer than one week unless directed by a doctor.

Ask a doctor before use if you

- have stomach pain, nausea or vomiting
- notice a change in bowel habits that last over two weeks
- are pregnant or breastfeeding

Stop use and ask a doctor if

you have rectal bleeding or failure to have a bowel movement after use.

Keep out of reach of children.

In case of accidental overdose, get medical help or contact a Poison Control Center immediately.

Directions

Shake well before using. Do not exceed maximum dose. Take at bedtime or as directed by a doctor.

Age	Usual Dose	Maximum Dose
Adults & children 12 years and older	2-3 teaspoons once a day	3 teaspoons twice a day

6 to 12 years	1 to 1-1/2 teaspoons once a	1 1/2 teaspoons twice a
	day	day
2 to 6 years	1/2 to 3/4 teaspoon once a day	3/4 teaspoon once a day
Under 2 years	Consult a doctor	

Other Information

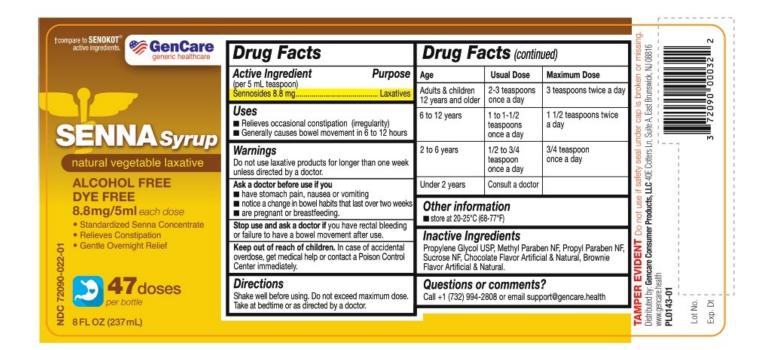
• store at 20-25°C (68-77°F)

Inactive Ingredients

Propylene Glycol USP, Methyl Paraben NF, Propyl Paraben NF, Sucrose NF, Chocolate Flavor Artificial & Natural, Brownie Flavor Artificial & Natural.

Questions or comments?

Call + 1 (732) 994-2808 or email support@gencare.health



SENNA SYRUP sennosides syrup					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72090-022		
Route of Administration	ORAL				
Active Ingredient/Active Moiety					

	Ingredient Name	Basis of Strength	Strength	
SENNOSIDES (UNII	: 3FYP5M0IJX) (SENNOSIDES - UNII:3FYP5M0IJX)	SENNOSIDES	8.8 mg in 5 mL	
Inactive Ingre	dients			
Ingredient Name			Strength	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
METHYLPARABEN (UNII: A2I8C7HI9T)				
PROPYLPARABEN (UNII: Z8IX2SC10H)				
SUCROSE (UNII: C1	51H8M554)			
Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:72090-022- 01	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/18/2024		
Marketing	Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Dru	ıg 505G(a)(3)	10/18/2024		
		10, 10, 20, 2021		

Labeler - Pioneer Life Sciences, LLC (014092742)

Registrant - Pioneer Life Sciences, LLC (014092742)

Revised: 10/2024

Pioneer Life Sciences, LLC