SENNA LAXATIVE- sennosides 8.6 mg tablet
NUVICARE LLC

Active Ingredient

Sennosides 8.6 mg

Purpose

Laxative

Uses:

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

Warnings:

Do not use

- Laxative products for longer than 1 week, unless directed by a doctor.

Ask a doctor before use if you have

- Stomach pain
- nausea
- vomiting,
- Noticed a sudden change in bowel habits that continues over a period of 2 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 (1-800-222-1222) right away.

DIRECTIONS

- Take preferably at bedtime or as directed by a doctor.

age	**starting dosage**	**maximum dosage**
adults & children 12 years of age and over	2 tablets once a day	4 tablets twice a day

children 6 to under 12 years	1 tablet once a day	2 tablets twice a day
children 2 to under 6 years	1 /2 tablet once a day	1 tablet twice a day
children under 2 years	ask a doctor	ask a doctor

OTHER INFORMATION

- store at 20-25°C (68-77°F) excursions permitted between15°-30°C (59°-86°F)
- each tablet contains: Calcium 25 mg & sodium 3 mg

INACTIVE INGREDIENTS

Colloidal silicon dioxide,croscarmellose sodium, Dicalcium phosphate, Hypromellose, liquid paraffin, magnesium stearate, microcrystalline cellulose, Maltodextrin, Purified water, Sodium lauryl sulphate, Stearic Acid.

QUESTIONS OR COMMENTS?

Call 1 (718) 337-8733 or visit: support@nuvicare.com

SENNA LAXATIVE

sennosides 8.6 mg tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84324-012
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SENNOSIDES (UNII: 3FYP5M0IJX) (SENNOSIDES - UNII:3FYP5M0IJX)	SENNOSIDES	8.6 mg

Inactive Ingredients

Ingredient Name	Strength
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID D7 (UNII: T3B081197X)	
PARAFFIN (UNII: I9O0E3H2ZE)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	

Product Characteristics

Color	brown	**Score**	no score
Shape	ROUND	**Size**	10mm
Flavor		**Imprint Code**	none
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84324-012-01	200 in 1 BOTTLE; Type 0: Not a Combination Product	07/17/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	07/17/2024	

Labeler - NUVICARE LLC (119257565)

Registrant - NUVICARE LLC (119257565)

Revised: 7/2024

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