SENNA NATURAL RELIEF- senna natural relief tablet
Pioneer Life Sciences, LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Active Ingredient

Sennosides 8.6 mg

Purpose

Laxative

Uses:

- relieves occasional constipation (irregularity).
- generally prodcues a bowel movement in 6-12 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

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 1-732-698-5070 Monday through Friday 9AM–5PM EST

PRINCIPAL DISPLAY PANEL

SENNA NATURAL RELIEF

senna natural relief tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72090-031
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:72090-031-60	600 in 1 BOTTLE; Type 0: Not a Combination Product	08/03/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	08/03/2020	

Labeler - Pioneer Life Sciences, LLC (014092742)