SENNA-LAX- sennosides tablet, film coated
SENNOSIDES- sennosides tablet, film coated
Major Pharmaceuticals

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.

1122 - Major

Drug Facts

Active ingredient (in each tablet)

Sennosides 8.6 mg

Purpose

Laxative

Uses

- relieves occasional constipation (irregularity)
- generally causes a bowel movement in 6-12 hours

Do not use

- laxative products for longer than one week unless directed by a doctor

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laxative products for longer than one 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 care 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

take preferably at bedtime or as directed by a doctor

age	starting dosage	maximum dosage

Adults and children 12 years of age and older	2 tablets once a day	4 tablets twice a day
Children 6 to under 12 years of age	1 tablet once a day	2 tablets twice a day
Children 2 to under 6 years of age	1/2 tablet once a day	1 tablet twice a day
Children under 2 years of age	ask a doctor	ask a doctor

Other information

- Each tablet contains: **Calcium 25 mg**
- Store at room temperature

Inactive ingredients

croscarmellose sodium, dicalcium phosphate, hypromellose, magnesium stearate, maltodextrin, microcrystalline cellulose, mineral oil, polyethylene glycol and talc

Questions or comments?

(800) 616-2471

Tamper Evident:

Do not use if sealed blister units are broken or damaged.

Product color may slightly vary

due to natural changes of ingredients.

Distributed by:

MAJOR® PHARMACEUTICALS

17177 N Laurel Park Drive, Suite 233

Livonia, MI 48152

NDC 0904-6725-80

MAJOR®

Senna Tablets

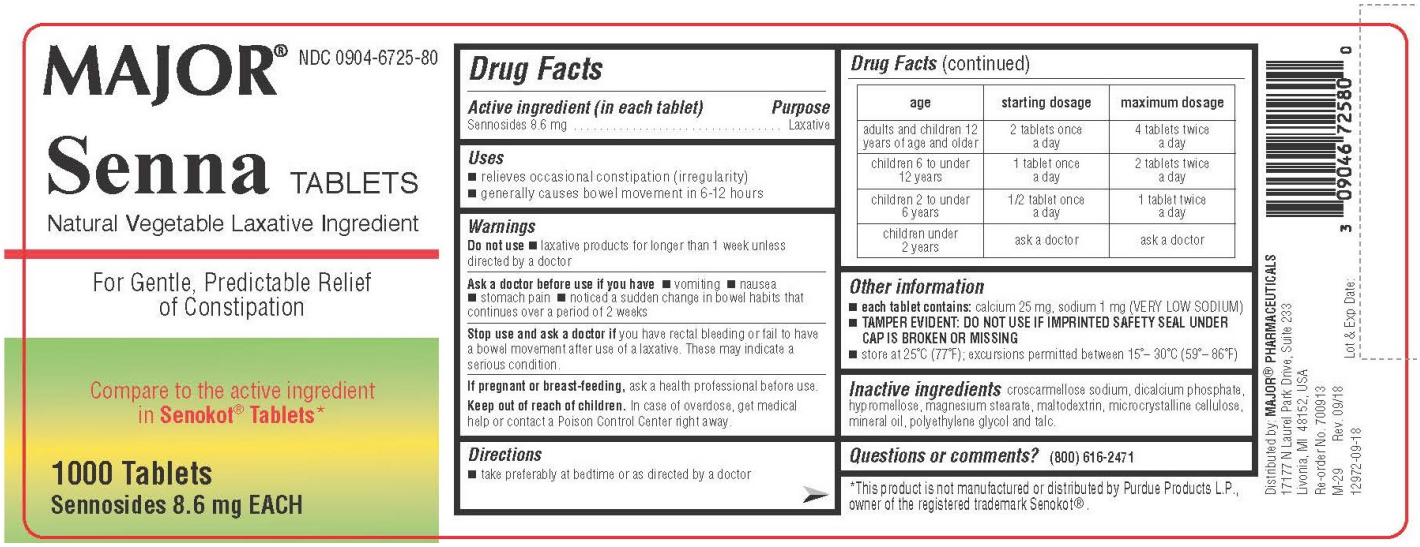
Natural Vegetable Laxative Ingredient

For Gentle, Predictable Relief of Constipation

Compare to the active ingredient in Senokot® Tablets*

Sennosides 8.6 mg EACH

1000 TABLETS



SENNA-LAX

sennosides tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0904-6522
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**
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
TALC (UNII: 7SEV7J4R1U)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	

Product Characteristics			
Color	brown	**Score**	no score
Shape	ROUND	**Size**	9mm
Flavor		**Imprint Code**	1122;1122
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0904-6522-61	10 in 1 BOX	03/07/2016	
1		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	03/07/2016	

SENNOSIDES

sennosides tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0904-6725
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
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
TALC (UNII: 7SEV7J4R1U)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	

Product Characteristics

Color	brown	**Score**	no score
Shape	ROUND	**Size**	9mm
Flavor		**Imprint Code**	1122;1122
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0904-6725-80	1000 in 1 BOTTLE; Type 0: Not a Combination Product	12/20/2018	
2	NDC:0904-6725-59	1 in 1 CARTON	12/20/2018	
2		100 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	12/20/2018	

Labeler - Major Pharmaceuticals (191427277)

Revised: 12/2020 Major Pharmaceuticals