# SENNA-LAX- sennosides tablet Preferred Pharmaceuticals Inc.

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.

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Senna-Lax

**Drug Facts** 

# Active ingredient

(in each tablet)

Sennosides 8.6 mg

## **Purpose**

Laxative

#### Uses

- relieves occasional constipation (irregularity)
- generally causes bowel movement in 6-12 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, 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**

- preferable at bedtime or a directed by a doctor
- adults and children 12 years of age and older: 2 tablets once a day: maximum of 4 tablets once a day
- children (6-12 years) 1tablet once a day: maximum 2 tablets once a day
- children (2-6 years) ½ tablet once a day: maximum 1 tablet once a day
- children under 2 years of age: Ask a doctor

### Other information

- Store at room temperature, USP
- Each tablet contains: Calcium 20 mg

# **Inactive ingredient**

Croscarmellose Sodium, Dicalcium Phosphate, Hypromellose, Magnesium Silicate, Magnesium Stearate, Microcrystalline Cellulose, Mineral Oil and Polyethylene Gycol

# 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

# Manufactured by:

Contract Pharmacal Corp. 135 Adams Avenue Hauppauge, NY 11788 USA www.cpc.com

# Distributed By:

## **MAJOR® PHARMACEUTICALS**

31778 Enterprise Drive

Livonia, MI 48150

Rev. 10/15

# Repackaged By: Preferred Pharmaceuticals Inc.

# **Principal Display Panel**

Senna-Lax

(Standardized Senna Concentrate)

8.6 mg Sennosides

100 Tablets

Senna-Lax Qty: Ins: Lot#: Bat#: Prod# (NDC): PREFERRED Senna-Lax CAUTION: Federal law PROHIBITS transfer of this drug to any person other than the patient for Generic for: Senokot Each tablet contains: Sennosides 8.6mg..Laxative whom it was prescribed. Pkg Size: Exp Date: Lot#: Batch#: Ins: Tomelo como se indica Scnna-Lax Qty: Ins: Lot#: Bat#: Prod# (NDC): Instrucciones Espanol: Mfg: Major Pharm.; Livonia, Ml Prod#: Directions English Proop

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# **SENNA-LAX**

sennosides tablet

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68788-6442(NDC:0904-6522)	
Route of Administration	ORAL			

Active Ingredient/Active Moiety

Ingredient Name
Basis of Strength
SENNOSIDES A AND B (UNII: 1B5FPI42EN) (SENNOSIDES A AND B - UNII:1B5FPI42EN)

SENNOSIDES A AND B 8.6 mg

Inactive Ingredients			
Ingredient Name	Strength		
CROSCARMELLOSE SODIUM (UNII: M28 OL1HH48)			
DIBASIC CALCIUM PHO SPHATE DIHYDRATE (UNII: O7TSZ97GEP)			
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)			
MAGNESIUM SILICATE (UNII: 9B9691B2N9)			
MAGNESIUM STEARATE (UNII: 70097M6130)			
MICRO CRYSTALLINE CELLULO SE (UNII: OP1R32D61U)			
LIGHT MINERAL OIL (UNII: N6K5787QVP)			
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)			

Product Characteristics			
Color	BROWN (Mottled brown)	Score	no score
Shape	ROUND	Size	9 mm
Flavor		Imprint Code	1122
Contains			

Packaging		
	Markating Start	Markating End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	3	30 in 1 BLISTER PACK; Type 0: Not a Combination Product	07/29/2016	
2	NDC:68788-6442-1	100 in 1 BLISTER PACK; Type 0: Not a Combination Product	07/29/2016	
3	NDC:68788-6442- 8	120 in 1 BLISTER PACK; Type 0: Not a Combination Product	06/23/2017	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	07/29/2016	

# Labeler - Preferred Pharmaceuticals Inc. (791119022)

# **Registrant - Preferred Pharmaceuticals Inc. (791119022)**

Establishment				
Name	Address	ID/FEI	Business Operations	
Preferred Pharmaceuticals Inc.		79 1119 0 22	REPACK(68788-6442)	

Revised: 6/2017 Preferred Pharmaceuticals Inc.