LAXACIN- docusate sodium and sennosides tablet
NuCare 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.

Laxacin

Laxative Plus Stool Softener

Active ingredient

Docusate Sodium 50 mg

Sennosides 8.6 mg

Purpose

Stool Softener

Laxative

Uses

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

Warnings

Do not use

- if you are now taking mineral oil, unless directed by a doctor
- 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 right away.

LOT#

Exp. Date:

Directions

- take preferably at bedtime or as directly by a doctor

age	starting dose	maximum dose
adults and children 12 years and older	2 tablets once a day	4 tablets twice a day

Other information

- Each tablet contains: Calcium 20 mg
- Each tablet contains: Sodium 4 mg
- Store at room temperature
- Keep lid tightly closed in a dry place
- Do not use if imprinted safety seal under cap is broken or missing

Inactive ingredients

Croscarmellose Sodium, D&C Yellow #10, Dextrose, Dicalcium Phosphate, FD&C Yellow #6, Hypromelose, Magnesium Stearate, Microcrystalline Cellulose, Polyethylene Glycol, Silica, Sodium Benzoate, Stearic Acid, Titanium Dioxide.

Questions? If you have any questions or comments, or to report an adverse event, please contact (480) 253-9761.

Manufactured for:
Alexso Inc.
Thousand Oaks, CA 91360

Package/Label Principal Display Panel

LAXACIN

docusate sodium and sennosides tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68071-1549(NDC:50488-0901)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DOCUSATE SODIUM (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG)	DOCUSATE SODIUM	50 mg
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: M28OL1HH48)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
DEXTROSE (UNII: IY9XDZ35W2)	
ANHYDROUS DIBASIC CALCIUM PHOSPHATE (UNII: L11K75P92J)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	orange	**Score**	no score
Shape	ROUND	**Size**	10mm
Flavor		**Imprint Code**	G55
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-1549-0	100 in 1 BOTTLE; Type 0: Not a Combination Product	12/19/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	10/01/2011	

Labeler - NuCare Pharmaceuticals,Inc. (010632300)

Establishment

Name	Address	ID/FEI	Business Operations
NuCare Pharmaceuticals,Inc.		010632300	relabel(68071-1549)