EX-LAX REGULAR STRENGTH STIMULANT LAXATIVE- sennosides pill
Novartis Consumer Health, 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.

Drug Facts

Active ingredient

Sennosides, USP, 15 mg

Purpose

Stimulant laxative

Uses

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

Warnings

Click here to enter Warnings

Do Not Use

Do not use laxative products when abdominal pain, nausea, or vomiting are present unless directed by a doctor

Ask Doctor before use if you have

noticed a sudden change in bowel habits that persists over a period of 2 weeks

Ask a doctor or pharmacist before use if you

are taking any other drug. Laxatives may affect how other drugs work. Take this product 2 or more hours before or after other drugs.

When Using this product

do not use for a period longer than 1 week

Stop use and ask a doctor if

rectal bleeding or failure to have a bowel movement occur after use of a laxative. These may be signs of 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.

Directions

- swallow pill(s) with a glass of water
- swallow pill(s) whole, do not crush, break or chew

adults and children 12 years of age and older	2 pills once or twice daily
children 6 to under 12 years of age	1 pill once or twice daily
children under 6 years of age	ask a doctor

Other information

- **each pill contains**: calcium 50 mg
- sodium free
- store at controlled room temperature 20-25C (68-77F). Protect from moisture.

Inactive ingredients

acacia, alginic acid, carnauba wax, colloidal silicon dioxide, dibasic calcium phosphate, iron oxides, magnesium stearate, microcrystalline cellulose, potassium hydroxide, pregelatinized starch, propylene glycol, shellac, sodium benzoate, sodium lauryl sulfate, stearic acid, sucrose, talc, titanium dioxide

Questions ?

call **1-800-452-0051**

Additional information listed on other panels

The Ex•Lax® Guarantee: When taken as directed, Ex•Lax® is guaranteed to work gently and effectively overnight or your money back. Return product to Novartis, attention Consumer Affairs, for full refund.

Tamper Evident Feature: Ex•Lax® Pills are sealed in blister packets. Use only if the individual seal is unbroken.

Distributed by: **Novartis Consumer Health, Inc.** Parsippany, NJ 07054-0622 ©20xx

Package/Label Principal Display Panel

NDC 0067-0003-30

Regular Strength

ex•lax®

SENNOSIDES, USP, 15 mg

STIMULANT LAXATIVE

RELIEF GUARANTEED EVERY TIME

GENTLE OVERNIGHT RELIEF YOU CAN TRUST

EX-LAX REGULAR STRENGTH STIMULANT LAXATIVE

sennosides pill

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0067-0003
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SENNOSIDES (UNII: 3FYP5M0IJX) (SENNOSIDES - UNII:3FYP5M0IJX)	SENNOSIDES	15 mg

Inactive Ingredients	
Ingredient Name	**Strength**
ACACIA (UNII: 5C5403N26O)	
ALGINIC ACID (UNII: 8C3Z4148WZ)	

CARNAUBA WAX (UNII: R12CBM0EIZ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CALCIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: L11K75P92J)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SHELLAC (UNII: 46N107B71O)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
SUCROSE (UNII: C151H8M554)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	WHITE (Beige)	**Score**	no score
Shape	ROUND	**Size**	11mm
Flavor		**Imprint Code**	ex;lax;1
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0067-0003-08	1 in 1 CARTON		
1		8 in 1 BLISTER PACK		
2	NDC:0067-0003-30	2 in 1 CARTON		
2		15 in 1 BLISTER PACK		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	01/01/2012	

Labeler - Novartis Consumer Health, Inc. (879821635)

Revised: 2/2013 Novartis Consumer Health, Inc.