EX-LAX REGULAR STRENGTH STIMULANT LAXATIVE- sennosides pill
EX-LAX REGULAR STRENGTH STIMULANT LAXATIVE- sennosides tablet
GlaxoSmithKline Consumer Healthcare Holdings (US) 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.

Drug Facts

Active ingredient (in each tablet)

Sennosides 15 mg

Purpose

Stimulant laxative

Uses

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

Warnings

Do not use

laxative products when abdominal pain, nausea, or vomiting are present unless directed by a doctor

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

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

Other information

- **each tablet contains**: calcium 45 mg, magnesium 5mg
- store at controlled room temperature 20-25°C (68-77°F)

Inactive ingredients (0067-0003)

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

Inactive Ingredients (0067-8141)

acacia, calcium carbonate, carnauba wax, corn starch, dibasic calcium phosphate, iron oxide black, iron oxide red, iron oxide yellow, magnesium stearate, methylparaben, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, propylene glycol, propylparaben, shellac, silicon dioxide, sodium benzoate, sodium lauryl sulfate, sucrose, talc, titanium dioxide

Questions ?

call **1-855-221-5432**

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

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.

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Distributed by: **Novartis Consuemr Health, Inc.**

Parsippany, NJ 07054-0622

Principal Display Panel

NDC 0067-8141-02

ex•lax®

SENNOSIDES, 15 mg

REGULAR STRENGTH

STIMULANT LAXATIVE

RELIEF GUARANTEED

GENTLE OVERNIGHT RELIEF YOU CAN TRUST

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 GSK, attention Consumer Affairs, for full refund.

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

Distributed by:

GSK Consumer Healthcare

Warren, NJ 07059

13143

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)	
ANHYDROUS DIBASIC CALCIUM PHOSPHATE (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	01/01/2012	11/30/2019
1		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:0067-0003-30	2 in 1 CARTON	01/01/2012	11/30/2019
2		15 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	01/01/2012	11/30/2019

EX-LAX REGULAR STRENGTH STIMULANT LAXATIVE

sennosides tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0067-8141
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)	
CALCIUM CARBONATE (UNII: H0G9379FGK)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
STARCH, CORN (UNII: O8232NY3SJ)	
ANHYDROUS DIBASIC CALCIUM PHOSPHATE (UNII: L11K75P92J)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL (UNII: 532B59J990)	
POVIDONES (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SHELLAC (UNII: 46N107B71O)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
SUCROSE (UNII: C151H8M554)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	BROWN (Tan)	**Score**	no score
Shape	ROUND	**Size**	10mm
Flavor		**Imprint Code**	ex;lax;1
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0067-8141-01	1 in 1 CARTON	07/01/2017	
1		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		

2	NDC:0067-8141-02	2 in 1 CARTON	07/01/2017	
2		15 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	07/01/2017	

Labeler - GlaxoSmithKline Consumer Healthcare Holdings (US) LLC (079944263)

Revised: 7/2016 GlaxoSmithKline Consumer Healthcare Holdings (US) LLC