EX-LAX MAXIMUM STRENGTH STIMULANT LAXATIVE- sennosides pill
Haleon US Holdings LLC

Drug Facts

Active ingredient (in each tablet)

Sennosides 25 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 40 mg, magnesium 5 mg
- store at controlled room temperature 20-25C (68-77F).

Inactive ingredients

acacia, calcium carbonate, carnauba wax, corn starch dibasic calcium phosphate, FD&C blue no.1 aluminum lake, iron oxide black, magnesium stearate, methyl paraben, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, propylene glycol, propylparaben, shellac, silicon dioxide, sodium benzoate, sodium lauryl sulfate, sucrose, talc, titanium dioxide

Questions or Comments?

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

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 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.

Principal Display Panel

MAXIMUM STRENGTH

ex•lax

SENNOSIDES, 25 mg

STIMULANT LAXATIVE

RELIEF GUARANTEED

24 TABLETS

GENTLE OVERNIGHT RELIEF YOU CAN TRUST

EX-LAX MAXIMUM STRENGTH STIMULANT LAXATIVE

sennosides pill

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0067-8142
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SENNOSIDES (UNII: 3FYP5M0IJX) (SENNOSIDES - UNII:3FYP5M0IJX)	SENNOSIDES	25 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)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
ALUMINUM OXIDE (UNII: LMI26O6933)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL 3350 (UNII: G2M7P15E5P)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
POVIDONE, UNSPECIFIED (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	BLUE	**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:0067-8142-01	2 in 1 CARTON	07/01/2017	
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:0067-8142-02	4 in 1 CARTON	07/01/2017	
2		12 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 Drug	M007	07/01/2017	

Labeler - Haleon US Holdings LLC (079944263)

Revised: 12/2024 Haleon US Holdings LLC