

LAXATIVE MAXIMUM STRENGTH- sennosides tablet, sugar coated
Chain Drug Consortium

Premier Value 44-348 Delisted

Active ingredient (in each tablet)

Sennosides USP, 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 (1-800-222-

1222) 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 and over	2 tablets once or twice daily
children 6 to under 12 years	1 tablet once or twice daily
children under 6 years	ask a doctor

Other information

- **each tablet contains:** calcium 40 mg, magnesium 5 mg
- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN**
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- see end flap for expiration date and lot number

Inactive ingredients

acacia, calcium carbonate, carnauba wax, corn starch, dibasic calcium phosphate dihydrate, FD&C blue #1 aluminum lake, iron oxide black, magnesium stearate, methylparaben, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, propylene glycol, propylparaben, shellac glaze, silicon dioxide, sodium benzoate, sodium lauryl sulfate, sucrose, talc, titanium dioxide

Questions or comments?

1-800-426-9391

Principal Display Panel

**Premier
Value®**

***COMPARE TO THE ACTIVE INGREDIENT IN
ex•lax® MAXIMUM STRENGTH**

Maximum Strength

Laxative

**Sennosides USP, 25 mg
STIMULANT LAXATIVE**

24 Tablets

actual
size

**Gentle, Dependable
Constipation Relief**

PV

INDEPENDENTLY TESTED
SATISFACTION GUARANTEED

**TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER
UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING**

*This product is not manufactured or distributed by
GSK Consumer Healthcare S.A., owner of the registered
trademark ex•lax® Maximum Strength.
50844 REV0419B34808

**Distributed By:
Pharmacy Value Alliance, LLC
407 East Lancaster Avenue,
Wayne, PA 19087**

**If for any reason you are
not satisfied with this product,
please return it to the store
where purchased
for a full refund.**

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Drug Facts
 COMPLETE PRODUCT INFORMATION
 KEEP OUTER PACKAGE FOR



COMPARE TO THE ACTIVE INGREDIENT IN
 ex•lax® MAXIMUM STRENGTH

Maximum Strength Laxative

Sennosides USP, 25 mg
 STIMULANT LAXATIVE

actual size



Gentle, Dependable
 Constipation Relief

24 Tablets



8 40986 03284 4
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B-1590-348-08
 REV/0419B34808

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 REV0419B34808
 50844
 Distributed by:
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 407 East Lancaster Avenue,
 Wayne, PA 19087

Drug Facts (continued)
 cellulose, polyethylene glycol, polyvinyl alcohol, povidone, propylene glycol, propylparaben, shellac glaze, silicon dioxide, sodium benzoate, sodium lauryl sulfate, sucrose, talc, titanium dioxide

Questions or comments? 1-800-426-9391

Drug Facts (continued)

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 ■ swallow tablet(s) with a glass of water
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adults and children	2 tablets once or	twice daily
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Inactive ingredients acacia, calcium carbonate, carnauba wax, corn starch, dibasic calcium phosphate, dithyrate, FD&C blue #1 aluminum lake, iron oxide black, magnesium stearate, methylparaben, microcrystalline

Drug Facts (continued)

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.
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Premier Value 44-348

LAXATIVE MAXIMUM STRENGTH

sennosides tablet, sugar coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68016-739
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: 5C5403N260)	
CALCIUM CARBONATE (UNII: H0G9379FGK)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
STARCH, CORN (UNII: O8232NY3SJ)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM)	
FERROSFERRIC OXIDE (UNII: XM0M87F357)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
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	44;348
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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1	NDC:68016-739-04	2 in 1 CARTON	01/02/2003	03/31/2026
1		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	505G(a)(3)		01/02/2003	03/31/2026

Labeler - Chain Drug Consortium (101668460)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	manufacture(68016-739) , pack(68016-739)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(68016-739) , pack(68016-739)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(68016-739)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(68016-739)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(68016-739)