

LAXATIVE- sennosides, 25 mg tablet, film coated
Granulation Technology, Inc.

Laxative Tablets
SENNOSIDES, 25 mg
STIMULANT LAXATIVE

Drug Facts

Active ingredient (in each tablet)

Sennosides, 25 mg

Purpose

Stimulant Laxative

Uses

- Relieves occasional constipation (irregularity)
- Generally produces a bowel movement in 6-12 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 two 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 two or more hours before or after other drugs.

When using this product do not use for a period longer than one 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.

Inactive ingredients

croscarmellose sodium, dibasic calcium phosphate dihydrate, microcrystalline cellulose, pregelatinized starch, colloidal silicon dioxide, magnesium stearate, polyvinyl alcohol, titanium dioxide, polyethylene glycol (PEG) 3350, talc, yellow iron oxide, FD&C blue 1 lake

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222).

Directions

- swallow tablet (s) with a glass of water.

- swallow tablet (s) whole, do not crush, break, or chew.

Age	Dosage
adults and children 12 years of age and over	2 tablets once or twice daily
children 6 to under 12 years	1 tablet once a day or twice daily
children under 6 years	Ask a doctor

Other information

- **Each tablet contains:** Calcium 40 mg, Magnesium 5 mg
- Store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- Use by expiration date on package

Questions or comments?

call **1-973-276-0740**

Principal Display Panel

NDC 63561-0186-09

Laxative Tablets Label

SENNOSIDES, 25 mg / STIMULANT LAXATIVE

Compare to ex.lax[®] Maximum Strength Active Ingredient

90 Film-Coated Tablets

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Drug Facts (continued under label)

MANUFACTURED BY:
DNB PRODUCTS
GRANULATION TECHNOLOGY, INC.
FAIRFIELD, NJ 07004
MANUFACTURED IN USA Rev. 06/25 LB0075

Discard Seal Empty & Replace Cap
PLASTIC BOTTLE

3 63561 01869 4

Peel here

Glue Area

STOP PEELING

Drug Facts (continued)

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NDC 63561-0186-02

Laxative Tablets Label

SENNOSIDES, 25 mg / STIMULANT LAXATIVE

Compare to ex.lax ® Maximum Strength Active Ingredient

1000 Film-Coated Tablets

NDC 63561-0186-02

Granulation Technology, Inc.

Dr. Nigalaye's Best



Laxative Tablets

SENNOSIDES 25 mg /
STIMULANT LAXATIVE

Maximum Strength

- Gentle, dependable constipation relief

1000 Film-Coated Tablets

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Drug Facts (continued under label)

MANUFACTURED BY:
DNB PRODUCTS
GRANULATION TECHNOLOGY, INC.
FAIRFIELD, NJ 07004
MANUFACTURED IN USA

Rev. 07/25 LB0088



Peel here

Glue Area

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LAXATIVE

sennosides, 25 mg tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63561-0186
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
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
STARCH, CORN (UNII: O8232NY3SJ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYVINYL ALCOHOL (UNII: 532B59J990)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POLYETHYLENE GLYCOL 3350 (UNII: G2M7P15E5P)	
TALC (UNII: 7SEV7J4R1U)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM)	

Product Characteristics

Color	blue	Score	no score
Shape	ROUND	Size	10mm
Flavor		Imprint Code	G186

Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63561-0186-9	90 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/11/2025	
2	NDC:63561-0186-2	1000 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/29/2025	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	505G(a)(3)	06/11/2025		

Labeler - Granulation Technology, Inc. (847132193)

Registrant - Granulation Technology, Inc. (847132193)

Establishment

Name	Address	ID/FEI	Business Operations
Granulation Technology, Inc.		847132193	manufacture(63561-0186)

Revised: 7/2025

Granulation Technology, Inc.