

SENNA LAXATIVE- sennosides tablet
Dolgenercorp, Inc. (DOLLAR GENERAL & REXALL)

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 8.6 mg

Purpose

Laxative

Uses

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

Warnings

Do not use

- laxative products for longer than 1 week unless directed by a doctor

Ask a doctor before use if you have

- stomach pain
- nausea
- vomiting
- noticed a sudden change in bowel habits that continues over a period of 2 weeks

Stop use and ask a doctor if

you have rectal bleeding or fail to have a bowel movement after use of a laxative. These may indicate 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

- take preferably at bedtime or as directed by a doctor

age	starting dosage	maximum dosage
adults and children 12 years of age or older	2 tablets once a day	4 tablets twice a day
children 6 to under 12 years	1 tablet once a day	2 tablets twice a day
children 2 to under 6 years	1/2 tablet once a day	1 tablet twice a day
children under 2 years	ask a doctor	ask a doctor

Other information

- **each tablet contains:** calcium 25 mg
- store at 25°C (77°F); excursions permitted between 15°C-30°C (59°F-86°F)

Inactive ingredients

croscarmellose sodium, dibasic calcium phosphate dihydrate, hypromellose*, liquid paraffin*, magnesium stearate, maltodextrin*, microcrystalline cellulose, mineral oil*, polyethylene glycol*, polyvinyl alcohol*, silicon dioxide*, stearic acid*, sodium lauryl sulfate*, talc*.

*contains one or more of these ingredients

Questions or comments?

Call **1-888-309-9030**

Principal Display Panel

COMPARE TO THE ACTIVE INGREDIENT IN SENOKOT®**

Senna Laxative

SENNOSIDES 8.6 mg

- Natural Vegetable Laxative Ingredient
- Gentle overnight relief

TABLETS

**This product is not manufactured or distributed by Purdue Products L.P., owner of the registered trademark Senokot®.

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

Package Label

DG health

Compare to the active
ingredient of Senokot[®]

Senna Laxative

Sennosides 8.6 mg/Laxative

Natural Vegetable Laxative Ingredient
Gentle, Overnight Relief

8.6
mg



100 Tablets

Actual Tablet Size

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY
SEAL UNDER CAP IS BROKEN OR MISSING.
KEEP OUTER CARTON FOR COMPLETE
WARNINGS AND PRODUCT INFORMATION.

100%
Satisfaction
Guaranteed!
(888) 309-9030

DISTRIBUTED BY DOLGENCORP, LLC
100 MISSION RIDGE
GOODLETTSVILLE, TN 37072

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Drug Facts (continued)

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children under 2 years	ask a doctor	ask a doctor

Directions

■ take preferably at bedtime or as directed by a doctor

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

If pregnant or breast-feeding, ask a health professional before use.

Indicate a serious condition.

Stop use and ask a doctor if you have rectal bleeding or fail to have a bowel movement after use of a laxative. These may

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

■ stomach pain ■ nausea ■ vomiting

Ask a doctor before use if you have

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Uses

Sennosides 8.6 mg.....Laxative

Active ingredient (in each tablet)

Purpose

Drug Facts

A0198



PLD-E22W FC002780

Lot No.:

Exp. Date:

Sennosides 8.6 mg laxative tablets

SENNALAXATIVE

sennosides tablet						
Product Information						
Product Type		HUMAN OTC DRUG	Item Code (Source)	NDC:55910-486		
Route of Administration		ORAL				
Active Ingredient/Active Moiety						
Ingredient Name			Basis of Strength	Strength		
SENNOSIDES A AND B (UNII: 1B5FPI42EN) (SENNOSIDES A AND B - UNII:1B5FPI42EN)			SENNOSIDES A AND B	8.6 mg		
Inactive Ingredients						
Ingredient Name				Strength		
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)						
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)						
HYPROMELLOSES (UNII: 3NXW29V3WO)						
PARAFFIN (UNII: I9O0E3H2ZE)						
MAGNESIUM STEARATE (UNII: 70097M6I30)						
MALTODEXTRIN (UNII: 7CVR7L4A2D)						
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)						
MINERAL OIL (UNII: T5L8T28FGP)						
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)						
STEARIC ACID (UNII: 4ELV7Z65AP)						
SODIUM LAURYL SULFATE (UNII: 368GB5141J)						
TALC (UNII: 7SEV7J4R1U)						
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)						
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)						
Product Characteristics						
Color	brown	Score	no score			
Shape	ROUND	Size	10mm			
Flavor		Imprint Code	0813;AV;TCL;080;S8			
Contains						
Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:55910-486-01	1 in 1 BOX	12/09/2015	04/26/2024		
1		100 in 1 BOTTLE; 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	12/09/2015	04/26/2024

Labeler - Dolgencorp, Inc. (DOLLAR GENERAL & REXALL) (068331990)

Registrant - P & L Development, LLC (079765031)

Revised: 10/2021

Dolgencorp, Inc. (DOLLAR GENERAL & REXALL)