VEGETABLE LAXATIVE- sennosides tablet, film coated
CHAIN DRUG MARKETING ASSOCIATION INC

Quality Choice 44-298

Active ingredient (in each tablet)

Sennosides USP, 8.6 mg

Purpose

Stimulant laxative

Uses

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

When using this product

do not exceed recommended dosage.

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 and over	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 30 mg, sodium 1 mg **Very Low Sodium**
- store at 25ºC (77ºF); excursions permitted between 15º-30ºC (59º-86ºF)
- protect from moisture
- see end flap for expiration date and lot number

Inactive ingredients

croscarmellose sodium, dibasic calcium phosphate dihydrate, hypromellose, magnesium stearate, maltodextrin, microcrystalline cellulose, polyethylene glycol, silicon dioxide

Questions or comments?

1-800-426-9391

Principal display panel

QC®
QUALITY
CHOICE

NDC 63868-579-01

Compare to the
Active Ingredient in
Senokot®*

Vegetable Laxative
Sennosides USP, 8.6 mg

Stimulant Laxative
Vegetable Laxative Ingredient
for Gentle Overnight Relief

100 Tablets

Actual Size

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

*This product is not manufactured or distributed by Avrio Health L.P., owner of the registered trademark Senokot®. 50844 REV1020C29812

100% QC SATISFACTION GUARANTEED

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Novi, MI 48375
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Questions: 800-935-2362

Quality Choice 44-298

VEGETABLE LAXATIVE

sennosides tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-579
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SENNOSIDES (UNII: 3FYP5M0IJX) (SENNOSIDES - UNII:3FYP5M0IJX)	SENNOSIDES	8.6 mg

Inactive Ingredients

Ingredient Name	Strength
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

Product Characteristics

Color	brown (light)	**Score**	no score
Shape	ROUND	**Size**	9mm
Flavor		**Imprint Code**	44;298
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-579-01	1 in 1 CARTON	12/08/2020	
1		100 in 1 BOTTLE, PLASTIC; 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)	12/08/2020	

Labeler - CHAIN DRUG MARKETING ASSOCIATION INC (011920774)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(63868-579)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(63868-579)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	pack(63868-579)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(63868-579)

Revised: 12/2023 CHAIN DRUG MARKETING ASSOCIATION INC