SENNA- sennosides tablet H E B

Drug Facts

Active ingredient (in each tablet)

Sennosides 8.6 mg

Purpose

Laxative

Uses

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

Warnings

Do not use

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

Ask a doctor before using 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 (1-800-222-1222) right away.

Directions

• take preferably at bedtime or as directed by a doctor

age	starting dosage	maximum dosage
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adults and children 12 years of age 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 tablets 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
- store at 25° C (77° F); excursions permitted between 15-30°C (59-86°F)

Inactive Ingredients

croscarmellose sodium, dibasic calcium phosphate dihydrate, hypromellose, magnesium stearate, microcrystalline cellulose, silicon dioxide, talc, triacetin

Questions or comments?

Call toll free 1-877-753-3935 Monday-Friday 9AM-5PM EST

Principal Display Panel

Compare to Senokot® active ingredient**

Senna Laxative

Sennosides 8.6 mg

Laxative

TABLETS

**This product is not manufactured or distributed by Aviro Health L.P., distributor of 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.

MADE WITH PRIDE AND CARE FOR H-E-B®, SAN ANTONIO, TX 78204

Product Label

Compare to Senokot® active ingredient NDC 37808-782-10 Senna Laxative Sennosides, 8.6 mg PLD-6732A Laxative FC007910 F actual 100 TABLETS size

MADE WITH PRIDE AND CARE FOR H-E-8®, SAN ANTONIO, TX 78204

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

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

This product is not manufactured or distributed by Avrio Health L.P., distributor of Senokot.

CINESTIONS OF COMMENTS? Call toll free: 1-877-753-3935 Monday-Friday 9AM-5PM EST

stearate, microcrystalline cellulose, silicon dioxide, talc, triacetin

Inactive ingredients anhydrous dibasic calcium phosphate, croscarmellose sodium, hypromellose, magnesium

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Dung Facts (continued)

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Do not use | laxative products for longer than 1 week unless directed by a doctor Warnings

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SƏSN

Laxative. Sennosides 8.6 mg. Purpose Active ingredient (in each tablet) Drug Facts

H-E-B Senna Laxative

sennosides tablet

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:37808-782

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CENNOCIDES (UNIV. REVDEMOUS) (CENNOCIDES LINIUS REVDEMOUS)	CENNOCIDEC	0 6

SENNOSIDES (UNII: 3FYP5M0IJX) (SENNOSIDES - UNII:3FYP5M0IJX) SENNOSIDES 8.6 mg

Inactive Ingredients

Ingredient Name	Strength
CROSCARMELLOSE SODIUM (LINII: M28011HH48)	

CROSCARMELLOSE SODIUM (UNII: M280L1HH48)

ANHYDROUS DIBASIC CALCIUM PHOSPHATE (UNII: L11K75P92J)

HYPROMELLOSES (UNII: 3NXW29V3WO)
MAGNESIUM STEARATE (UNII: 70097M6I30)

CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)

SILICON DIOXIDE (UNII: ETJ7Z6XBU4)

TALC (UNII: 7SEV7J4R1U)

TRIACETIN (UNII: XHX3C3X673)

Product Characteristics

Color	brown	Score	no score
Shape	ROUND	Size	9mm
Flavor		Imprint Code	PS23

Contains

Packaging

# Item Code	Package Description	Marketing Start Date	Marketing End Date		
1 NDC:37808- 782-10	1 in 1 BOX	02/18/2022			
1	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product				

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
OTC Monograph Drug	505G(a)(3)	02/18/2022	

Revised: 8/2024 H E B