STOOL SOFTENER AND STIMULANT LAXATIVE- docusate sodium, sennosides tablet

H-E-B

HEB 209R Sennosides 8.6mg / Docusate Sodium 50mg 240ct

Drug Facts

Active ingredients (in each tablet)

Docusate sodium 50 mg

Sennosides 8.6 mg

Purposes

Stool Softener

Stimulant laxative

USES

Uses

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

Warnings

Do not use

- laxative products for longer than 1 week unless told to do so by a doctor
- if you are presently taking mineral oil unless told to do so by a doctor

Ask a doctor before use if you have

- stomach pain
- nausea '
- vomiting
- noticed a sudden change in bowel habits that lasts over 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 could 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 right away (1-800-222-1222).

Directions

• take only by mouth. Doses may be taken as a single daily dose, preferably in the evening, or in divided doses.

under 12 years of age children 2 to under 6 years of age children under 2

take 1-2 tablets daily

take up to 1 tablet daily

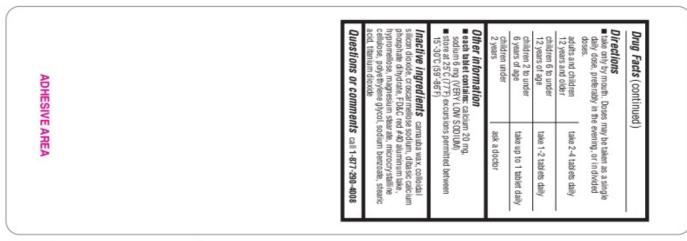
ask a doctor

Other information

- each tablet contains: calcium 20 mg, sodium 6 mg (VERY LOW SODIUM)
- Store at 25°C (77°F) excursions permitted between 15°-30°C (59°-86°F)

Inactive ingredients carnauba wax, colloidal silicon dioxide, croscarmellose sodium, dibasic calcium phosphate dihydrate, FD&C red #40 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, sodium benzoate, stearic acid, titanium dioxide

Questions or comments call 1-877-290-4008





STOOL SOFTENER AND STIMULANT LAXATIVE docusate sodium, sennosides tablet Product Information Product Type HUMAN OTC DRUG Item Code (Source) Route of Administration ORAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
DOCUSATE SODIUM (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG)	DOCUSATE SODIUM	50 mg		
SENNOSIDES (UNII: 3FYP5M0IJX) (SENNOSIDES - UNII:3FYP5M0IJX)	SENNOSIDES	8.6 mg		

Inactive Ingredients			
Ingredient Name	Strength		
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			
HYPROMELLOSES (UNII: 3NXW29V3WO)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
CARNAUBA WAX (UNII: R12CBM0EIZ)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
SODIUM BENZOATE (UNII: OJ245FE5EU)			
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)			
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)			
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)			
FD&C RED NO. 40 ALUMINUM LAKE (UNII: 6T47AS764T)			
STEARIC ACID (UNII: 4ELV7Z65AP)			

Product Characteristics				
Color	red	Score	no score	
Shape	ROUND	Size	10mm	
Flavor		Imprint Code	TCL097	
Contains				

Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:37808-549- 29	240 in 1 BOTTLE; Type 0: Not a Combination Product	05/04/2022	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	02/04/2022	

Labeler - H-E-B (007924756)

Registrant - TIME CAP LABORATORIES, INC. (037052099)

Establishment				
Name	Address	ID/FEI	Business Operations	
TIME CAP LABORATORIES, INC.		037052099	manufacture(37808-549)	

Revised: 2/2025 H-E-B