

**STOOL SOFTENER AND STIMULANT LAXATIVE- docusate sodium, sennosides tablet**  
**H-E-B**

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**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.

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adults and children  
12 years and older  
children 6 to

take 2-4 tablets daily

ask a doctor

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37808-549
Route of Administration	ORAL		

**Active Ingredient/Active Moiety**

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

**Inactive Ingredients**

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

**Product Characteristics**

<b>Color</b>	red	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	10mm
<b>Flavor</b>		<b>Imprint Code</b>	TCL097
<b>Contains</b>			

**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)