STOOL SOFTENER PLUS STIMULANT LAXATIVE- docusate sodium and sennosides tablet, film coated Meijer, Inc.

GS 303B (304)

Active ingredient (in each tablet)

Docusate Sodium 50 mg Sennosides 8.6 mg

Purpose

Stool softener

Stimulant laxative

Uses

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

Warnings

Do not use

- if you are now taking mineral oil, unless told to do so by a doctor
- 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 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 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 only by mouth. Doses may be taken as a single daily dose, preferably in the evening, or in divided doses

adults and children 12	take 2-4
years and over	tablets daily
children under 12 years	ask a doctor

Other information

- each tablet contains: calcium 20 mg, sodium 3 mg
- store at controlled room temperature
- **Tamper Evident:** Do not use if imprinted seal under cap is missing or broken.
- product of India

Inactive ingredients

cellulose, croscarmellose sodium, dicalcium phosphate, FD&C blue #2 lake, FD&C red #40 lake, hypromellose, magnesium stearate, PEG, silica, talc, titanium dioxide.

Questions or comments?

1-800-540-3765

Package Label



STOOL SOFTENER PLUS STIMULANT LAXATIVE

docusate sodium and sennosides tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79481-0304
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		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
HYPROMELLOSES (UNII: 3NXW29V3WO)			
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)			
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)			
CALCIUM PHOSPHATE, DIBASIC, DIHYDRATE (UNII: O7TSZ97GEP)			
FD&C RED NO. 40 (UNII: WZB9127XOA)			
SILICON DIOXIDE (UNII: ETJ7Z 6XBU4)			
TALC (UNII: 7SEV7J4R1U)			

Product Characteristics				
Color	red	Score	no score	
Shape	ROUND	Size	10mm	
Flavor		Imprint Code	PSD21	
Contains				

l	Packaging			
# Item Code Package Description		Marketing Start Date	Marketing End Date	
	1 NDC:79481-0304-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2025	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	03/01/2025	

Labeler - Meijer, Inc. (006959555)

Registrant - Geri-Care Pharmaceutical Corp (611196254)

Revised: 3/2025 Meijer, Inc.