STOOL SOFTENER PLUS STIMULANT LAXATIVE- docusate sodium and sennosides tablet McKesson (Sunmark)

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

DRUG FACTS

Active Ingredients (in each tablet)

Docusate Sodium 50 mg Sennosides 8.6 mg

Purpose

Stool softener Stimulant Laxative

Uses

- for overnight relief from 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 breastfeeding,

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.

Other information

- each tablet contains: calcium 20 mg
- each tablet contains: sodium 6 mg VERY LOW SODIUM
- store at 15° 30° C (59°-86° F), protect from excessive moisture

Inactive Ingredients

carnauba wax*, colloidal silicon dioxide, croscarmellose sodium, dibasic calcium phosphate dihydrate, FD&C red #40 aluminum lake,

FD&C yellow #6 aluminum lake*, hypromellose, magnesium stearate, maltodextrin*, microcrystalline cellulose, polydextrose*, polyethylene glycol*, purified water*, sodium benzoate*, stearic acid,talc*, titanium dioxide, triglycerides*

Questions or comments?

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

Principal Display Panel

**Compare to Peri- colace® active ingredients

**This product is not manufactured or distributed by Purdue products L.P., owner of the registered trademark Peri-Colace®

Stool Softener plus stimulant laxative

For relief of occasional constipation

Docusate Sodium 50 mg Sennosides 8.6 mg

Another Quality Product Distributed by McKesson

One post street, San Francisco, CA 94104

Money Back Guarantee

Please visit us at www.sunmarkbrand.com

Tamper Evident: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

Product Label

^{*}contains one or more of these ingredients





Stool softener plus stimulant laxative

STOOL SOFTENER PLUS STIMULANT LAXATIVE

docusate sodium and sennosides tablet

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49348-544	
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
SENNO SIDES (UNII: 3FYP5M0 IJX) (SENNO SIDES - UNII:3FYP5M0 IJX)	SENNOSIDES	8.6 mg

ı	Inactive Ingredients	
	Ingredient Name	Strength

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CARNAUBA WAX (UNII: R12CBM0 EIZ)	
CROSCARMELLOSE SODIUM (UNII: M28 OL 1 HH48)	
CALCIUM PHO SPHATE, DIBASIC, DIHYDRATE (UNII: O7TSZ97GEP)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTO DEXTRIN (UNII: 7CVR7L4A2D)	
CELLULOSE, MICRO CRYSTALLINE (UNII: OP1R32D61U)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)	
C10-18 TRIGLYCERIDES (UNII: 43AGM4PHPI)	
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)	
TALC (UNII: 7SEV7J4R1U)	
ALUMINUM OXIDE (UNII: LMI26O6933)	

Product Characteristics				
Color	RED	Score	no score	
Shape	ROUND	Size	10 mm	
Flavor		Imprint Code	TCL97;SS2;S44	
Contains				

ı	Packaging					
	# Item Code	Package Description	Marketing Start Date	Marketing End Date		
	1 NDC:49348-544-19	250 in 1 BOTTLE; Type 0: Not a Combination Product	07/09/2010	07/09/2020		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part334	07/09/2010	07/09/2020

Labeler - McKesson (Sunmark) (177667227)

Revised: 6/2019 McKesson (Sunmark)