

**STOOL SOFTENER PLUS STIMULANT LAXATIVE- docusate sodium and sennosides tablet  
MEIJER, INC.**

*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.*

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**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 directed by a doctor
- if you are now taking mineral oil, 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 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 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.

### 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 years and older	take 2-4 tablets daily
children 6 to under 12 years of age	take 1-2 tablets daily
children 2 to under 6 years of age	take up to 1 tablet daily
children under 2	ask a doctor

### Other information

- each tablet contains: **calcium 20 mg**
- each tablet contains: **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\*, croscarmellose sodium, dibasic calcium phosphate dihydrate, FD&C blue #2 aluminum lake, FD&C red #40 aluminum lake, hypromellose, magnesium stearate, maltodextrin\*, microcrystalline cellulose, polyethylene glycol\*, polyvinyl alcohol\*, silicon dioxide, sodium benzoate\*, stearic acid, talc\*, titanium dioxide

\*contains one or more of these ingredients

### Questions or comments?

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

### Principal Display Panel

Compare to Peri-Colace® active ingredients\*\*

Overnight Relief

**stool softener**

**Plus Stimulant Laxative**

Docosate Sodium, 50 mg / Sennosides, 8.6 mg

Relief of Occasional Constipation

- Effective
- Reliable
- Comfortable

TABLETS

**TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS**

**BROKEN OR MISSING.**

\*\*This product is not manufactured or distributed by Purdue Products LP., distributor of Peri-Colace®.

DIST. BY MEIJER DISTRIBUTION, INC.

GRAND RAPIDS, MI 49544

www.meijer.com

**Product Label**

NDC 41250-084-01 Compare To Peri-Colace® active ingredients\*\*



**Overnight Relief**

**Stool Softener Plus Stimulant Laxative**

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100 Tablets



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**Drug Facts (continued under label)**

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PLD-F9T  
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**Drug Facts (continued)**

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**Meijer Stool Softener Plus Stimulant Laxative**

**STOOL SOFTENER PLUS STIMULANT LAXATIVE**

docosate sodium and sennosides tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41250-084
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>CARNAUBA WAX</b> (UNII: R12CBM0EIZ)	
<b>CROSCARMELOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>DIBASIC CALCIUM PHOSPHATE DIHYDRATE</b> (UNII: O7TSZ97GEP)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>HYPROMELLOSES</b> (UNII: 3NXW29V3WO)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>MALTODEXTRIN</b> (UNII: 7CVR7L4A2D)	
<b>CELLULOSE, MICROCRYSTALLINE</b> (UNII: OP1R32D61U)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ05DW1A)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>ALUMINUM OXIDE</b> (UNII: LMI26O6933)	
<b>FD&amp;C BLUE NO. 2</b> (UNII: L06K8R7DQK)	
<b>POLYVINYL ALCOHOL, UNSPECIFIED</b> (UNII: 532B59J990)	

## 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;0806;AV;S44
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41250-084-01	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/30/2015	12/31/2024

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	11/30/2015	12/31/2024

