

STOOL SOFTENER WITH STIMULANT LAXATIVE- docusate sodium and sennosides tablet

P & L Development, LLC

Drug Facts

Active ingredients (in each tablet)

Docusate sodium 50 mg

Sennosides 8.6 mg

Purpose

Stool softener

Stimulant laxative

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 directed 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 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 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 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 21 mg
- **each tablet contains:** sodium 5 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 red #40 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, silicon dioxide, sodium benzoate, stearic acid, talc, titanium dioxide

Questions or comments?

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

Principal Display Panel

stool softener + stimulant laxative

docusate sodium 50 mg

(stool softener)

sennosides 8.6 mg

(stimulant laxative)

tablets

*Compare to the active ingredients in Colace® 2-IN-1

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

*This product is not manufactured or distributed by Avrio Health LP., distributor of Colace® 2-IN-1

Distributed by: **PL Developments**

200 Hicks Street, Westbury, NY 11590

Product Label

wb
wellness basics

stool softener
+ stimulant laxative
docusate sodium 50 mg
(stool softener)
sennosides 8.6 mg
(stimulant laxative)

100 tablets

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in Colace® 2-IN-1

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Drug Facts (continued under label)

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Distributed by: **PL Developments**
200 Hicks Street, Westbury, NY 11590

Actual Size

PLD-0677A Lot No.: 3 59726 48410 7 Exp. Date:
LB007897

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Drug Facts (continued)

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WELLNESS BASICS Stool Softener

STOOL SOFTENER WITH STIMULANT LAXATIVE

docusate sodium and sennosides tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59726-878
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
CARNAUBA WAX (UNII: R12CBM0EIZ)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
ALUMINUM OXIDE (UNII: LMI26O6933)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	

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:59726-878-40	400 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/29/2020	01/01/2027
2	NDC:59726-878-10	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/29/2020	01/01/2027

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	05/29/2020	01/01/2027

Labeler - P & L Development, LLC (800014821)