

**STOOL SOFTENER LAXATIVE- docusate sodium capsule, liquid filled**  
**P & L Development, LLC**

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**Drug Facts**

**Active ingredient (in each softgel)**

Docusate sodium 100 mg

**Purpose**

Stool softener laxative

**Uses**

- relieves occasional constipation (irregularity)
- generally produces bowel movement in 12 to 72 hours

**Warnings**

**Do not use**

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.
- you need to use a laxative for more than 1 week

**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 (1-800-222-1222) right away.

**Directions**

- take only by mouth. Doses may be taken as a single daily dose or in divided doses.

adults and children 12 years and over	take 1-3 softgels daily
children 2 to under 12 years of age	take 1 softgel daily
children under 2 years	ask a doctor

### Other information

- **each softgel contains:**sodium 6 mg
- store at 25°C (77°F); excursions permitted between 15-30°C (59-86°F)

### Inactive ingredients

black iron oxide, D&C red #33, FD&C blue #1, FD&C red #40, FD&C yellow #6, gelatin, glycerin, hypromellose, polyethylene glycol, propylene glycol, purified water, sorbitan, sorbitol, titanium dioxide

### Questions or comments?

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

### Principal Display Panel

†Compare to the active ingredient in Colace® Regular Strength Stool Softener

#### **stool softener**

docusate sodium 100 mg

stool softener laxative

Softgels

†This product is not manufactured or distributed by Avrio Health L.P., distributor of Colace Regular Strength Stool Softener.

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

Distributed by: **PL Developments**

200 Hicks Street, Westbury, NY 11590

### Product Label

**stool softener**  
docusate sodium  
100 mg  
stool softener laxative  
100 softgels

**wb**  
wellness basics

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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  
3 59726 23801 4  
PLD-8717A Lot No.: Exp. Date:  
LB008214

**PEEL HERE** →

**Drug Facts (continued)**  
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**WELLNESS BASICS Stool Softener Laxative**

**STOOL SOFTENER LAXATIVE**

docusate sodium capsule, liquid filled

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:59726-853
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>DOCUSATE SODIUM</b> (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG)	DOCUSATE SODIUM	100 mg

**Inactive Ingredients**

Ingredient Name	Strength
<b>D&amp;C RED NO. 33</b> (UNII: 9DBA0SBB0L)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>FD&amp;C YELLOW NO. 6</b> (UNII: H77VEI93A8)	
<b>GELATIN</b> (UNII: 2G86QN327L)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	

<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SORBITAN</b> (UNII: 6O92ICV9RU)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>FERROSO FERRIC OXIDE</b> (UNII: XM0M87F357)	

### Product Characteristics

<b>Color</b>	red, white	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	12mm
<b>Flavor</b>		<b>Imprint Code</b>	P10
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59726-853-10	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/31/2020	02/01/2027
2	NDC:59726-853-01	1000 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/31/2020	02/01/2027
3	NDC:59726-853-40	400 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/31/2020	02/01/2027

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	12/31/2020	02/01/2027

**Labeler** - P & L Development, LLC (800014821)

Revised: 11/2025

P & L Development, LLC