

STOOL SOFTENER EXTRA STRENGTH- docusate sodium capsule, liquid filled P & L Development, LLC

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 ingredient (in each softgel)

Docusate sodium 250 mg

Purpose

Stool softener laxative

Uses

- for relief of occasional constipation
- this product generally produces a bowel movement within 12 to 72 hours

Warnings

Do not use

if you are presently 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 last 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 right away.

Directions

- adults and children 12 years of age and over: take 1 softgel daily or as directed by a doctor
- children under 12 years of age: ask a doctor

Other information

- **each softgel contains:** sodium 15 mg
- store between 20-25°C(68-77°F); excursions permitted between 15-30°C (59-86°F)

Inactive Ingredients

edible white ink, FD&C red #40, FD&C yellow #6, gelatin, glycerin, mannitol, polyethylene glycol, propylene glycol, purified water, sorbitan, sorbitol

Questions or comments?

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

Principal Display Panel

extra strength

stool softener

docusate sodium 250 mg

stool softener laxative

- Stimulant-free
- relief of constipation

softgels

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



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

Drug Facts	Purpose Stool softener laxative
Active ingredient (in each softgel) Docusate sodium 250 mg	
Uses <ul style="list-style-type: none"> for relief of occasional constipation this product generally produces a bowel movement within 12 to 72 hours 	
Warnings Do not use if you are presently taking mineral oil, unless directed by a doctor. Ask a doctor before use if you have <ul style="list-style-type: none"> stomach pain nausea vomiting noticed a sudden change in bowel habits that lasts over 2 weeks Stop use and ask a doctor if <ul style="list-style-type: none"> 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 right away.	
Directions <ul style="list-style-type: none"> adults and children 12 years of age and over: take 1 softgel daily or as directed by a doctor children under 12 years of age: ask a doctor 	
Other information <ul style="list-style-type: none"> each softgel contains: sodium 15 mg store between 20-25°C (68-77°F); excursions permitted between 15-30°C (59-86°F) 	
Inactive ingredients edible white ink, FD&C red #40, FD&C yellow #6, gelatin, glycerin, mannitol, polyethylene glycol, propylene glycol, purified water, sorbitan, sorbitol	
Questions or comments? Call 1-877-753-3835 Monday-Friday 9AM-5PM EST	

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



PLD-B610A
LB007149

Lot No.:

Exp. Date:

WELLNESS BASICS Extra Strength Stool Softener

STOOL SOFTENER EXTRA STRENGTH

docusate sodium capsule, liquid filled

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59726-179
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DOCUSATE SODIUM (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG)	DOCUSATE SODIUM	250 mg

Inactive Ingredients

Ingredient Name	Strength
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
SORBITAN (UNII: 6O92ICV9RU)	
MANNITOL (UNII: 3OWL53L36A)	

Product Characteristics

Color	orange	Score	no score
--------------	--------	--------------	----------

Shape	CAPSULE	Size	20mm
Flavor		Imprint Code	P20
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59726-179-25	250 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/29/2019	11/29/2025

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	11/29/2019	11/29/2025

Labeler - P & L Development, LLC (800014821)

Revised: 4/2023

P & L Development, LLC