

**STOOL SOFTENER LAXATIVE EXTRA STRENGTH- docusate sodium capsule,
liquid filled**

P & L Development, LLC

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 13 mg
- store between 20-25°C(68-77°F); excursions permitted between 15-30°C (59-86°F)

Inactive Ingredients

anhydrous citric acid, FD&C red #40, FD&C yellow #6, gelatin, glycerin, isopropyl alcohol, lecithin, mannitol, mineral oil, polyethylene glycol, propylene glycol, purified water, sorbitan, sorbitol, white ink

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

relieves occasional constipation

Stimulant-free

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



NDC 59726-059-10

extra strength stool softener

docusate sodium 250 mg stool softener laxative

relieves occasional constipation stimulant-free

100 softgels



actual size

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

Drug Facts
Active ingredient (in each softgel) Docusate sodium 250 mg.....Stool softener laxative
Uses <ul style="list-style-type: none"> relieves occasional constipation (irregularity) this product generally produces a bowel movement within 12 to 72 hours
Warnings <p>Do not use if you are presently taking mineral oil, unless directed by a doctor.</p> <p>Ask a doctor before use if you have</p> <ul style="list-style-type: none"> stomach pain nausea vomiting <p>noticed a sudden change in bowel habits that lasts over 2 weeks</p> <p>Stop use and ask a doctor if</p> <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 <p>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.</p>
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 13 mg store between 20-25°C (68-77°F); excursions permitted between 15-30°C (59-86°F)
Inactive ingredients anhydrous citric acid, FD&C red #40, FD&C yellow #6, gelatin, glycerin, isopropyl alcohol, lecithin, mannitol, mineral oil, polyethylene glycol, propylene glycol, purified water, sorbitan, sorbitol, white ink
Questions or comments? Call 1-877-753-3935 Monday-Friday 9AM-5PM EST

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



3 597261285100
PLD-F727B Lot No.:
LB010763 Exp. Date:

READYinCASE Extra Strength Stool Softener

STOOL SOFTENER LAXATIVE EXTRA STRENGTH

docusate sodium capsule, liquid filled

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59726-059
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: 3WJQ05DW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
MANNITOL (UNII: 3OWL53L36A)	
MINERAL OIL (UNII: T5L8T28FGP)	
SORBITAN (UNII: 6O921CV9RU)	

Product Characteristics

Color	orange	Score	no score
Shape	CAPSULE	Size	20mm
Flavor		Imprint Code	P4
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59726-059-10	100 in 1 BOTTLE; Type 0: Not a Combination Product	04/16/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	04/16/2021	

Labeler - P & L Development, LLC (800014821)

Revised: 1/2026

P & L Development, LLC