

**FLEET- docusate sodium capsule, gelatin coated**  
**C.B. Fleet Company, 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 Ingredient**

(in each softgel)

Docusate Sodium 100 mg.....Stool Softener

**Uses**

- for the prevention of dry, hard stools
- for relief of occasional constipation

**If pregnant or breast-feeding**

Ask a health care professional before use.

**Keep out of reach of children**

In case of overdose, get medical help or contact a Poison Control Center right away.

**Do not use**

- if you are currently taking mineral oil, unless directed by a doctor
- when abdominal pain, nausea, or vomiting are present
- for longer than one week, unless directed by a doctor

**Ask a doctor before use if**

- you notice a sudden change in bowel habits that persists over a period of 2 weeks

**Stop use and ask a doctor if**

- you have rectal bleeding
- you fail to have a bowel movement after use

These symptoms may indicate a serious condition.

**Directions**

Single Daily Dosage

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

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## Other Information

- each softgel contains: sodium 5 mg
- this product generally produces a bowel movement within 12 to 72 hours
- mouth of bottle sealed for your safety. If foil imprinted "SEALED FOR YOUR PROTECTION" is broken or missing, do not use

## Inactive Ingredients

edible white ink, FDandC red 40, FDandC yellow 6, gelatin, glycerin, mannitol, polyethylene glycol, polysorbate, propylene glycol, purified water and sorbitol

## Questions?

1-866-255-6960 or [www.fleetlabs.com](http://www.fleetlabs.com)



**Fleet**<sup>®</sup>  
laxative

Sof-Lax<sup>®</sup>  
gentle  
stool softener

prevents constipation  
eases discomfort  
of hard stools



**#1 DOCTOR  
RECOMMENDED  
LAXATIVE BRAND** | **60** Soft Gels

## FLEET

docusate sodium capsule, gelatin coated

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0132-0751
Route of Administration	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
DOCUSATE SODIUM (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG)	DOCUSATE SODIUM	100 mg in 60

**Inactive Ingredients**

Ingredient Name	Strength
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
MANNITOL (UNII: 3OWL53L36A)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0K00R)	
SORBITOL (UNII: 506T60A25R)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	

**Product Characteristics**

Color	red (Clear, Bright Red)	Score	no score
Shape	OVAL	Size	5mm
Flavor		Imprint Code	P51
Contains			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0132-0751-60	60 in 1 BOTTLE, PLASTIC		

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	06/30/2002	

**Labeler** - C.B. Fleet Company, Inc. (003119054)

Revised: 7/2014

C.B. Fleet Company, Inc.