

**STOOL SOFTENER EXTRA STRENGTH- docusate sodium capsule, liquid filled
Mckesson (Sunmark)**

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

Uses

- for the prevention of dry, hard stools
- for relief of occasional constipation.
- this product generally produces a bowel movement within 12 to 72 hours.

Warnings - Do not use

- if you are taking mineral oil, unless directed by a doctor
- when abdominal pain, nausea, or vomiting are present
- for more than 1 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

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

Other information

- **each softgel contains:** sodium 15 mg
- store at controlled room temperature 15° - 30° C (59°- 86° F)
- do not use if imprinted safety seal under cap is broken or missing

Inactive Ingredients

edible white ink, FD&C Red No# 40, FD&C Yellow No# 6, gelatin, glycerin, polyethylene glycol, propylene glycol, purified water and sorbitol special.

Questions or comments?

Call toll free 1-877-753-3935

Principal Display Panel

Stool Softener

Extra strength

Relieves constipation

Docusate sodium 250 mg

Another quality product distributed by McKesson

one post street

San Francisco CA 94104

Money back gurantee

Please visit us at www.sunmarkbrand.com

Product Label

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Active ingredient (in each softgel) Docusate Sodium 250 mg..... Stool softener

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

Uses ■ for the prevention of dry, hard stools ■ for relief of occasional constipation ■ this product generally produces a bowel movement within 12 to 72 hours

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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

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: take as directed by a doctor

Other information

■ each softgel contains: sodium 15 mg

■ store at controlled room temperature 15°-30°C (59°-86°F)

■ do not use if imprinted safety seal under cap is broken or missing

Inactive ingredients edible white ink, FD&C Red #40, FD&C Yellow #6, gelatin, glycerin, polyethylene glycol, propylene glycol, purified water, and sorbitol special.

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Empowering Healthcare

Another Quality Product
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PLD-B
L15SMT00

Lot No.:
Exp. Date:

0 110939 98133 2

Docusate Sodium 250 mg

STOOL SOFTENER EXTRA STRENGTH

docusate sodium capsule, liquid filled

Product Information

Product Type	HUMAN OTC DRUG LABEL	Item Code (Source)	NDC:49348-714
Route of Administration	ORAL	DEA Schedule	

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DOCUSATE SODIUM (DOCUSATE)	DOCUSATE SODIUM	250 mg

Inactive Ingredients

Ingredient Name	Strength
FD&C RED NO. 40	
FD&C YELLOW NO. 6	
GELATIN	
GLYCERIN	
POLYETHYLENE GLYCOLS	
PROPYLENE GLYCOL	
WATER	
SORBITOL	

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:49348-714-10	100 in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part334	07/09/2010	

Labeler - Mckesson (Sunmark) (177667227)

Revised: 9/2012

Mckesson (Sunmark)