STOOL SOFTENER- docusate sodium capsule
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 100 mg

Purpose

Stool softener

Uses

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

Warnings - Do not use

- if abdominal pain, nausea or vomiting are present

Ask a doctor before use if

- you notice a sudden change in bowel habits that persists over a period of 2 weeks
- you are presently taking mineral oil

Stop use and ask a doctor if

- rectal bleeding or failure to have a bowel movement occur after use which may indicate a serious condition
- you need to use a laxative for more than 1 week

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: **1-800-222-1222**

Directions

- take with a glass of water

adults and children 12 years of age and over	take 1 to 3 softgels daily. This dose may be taken as a single daily dose or in divided doses.
children 2 to under 12 years of age	take 1 softgel daily
children under 2 years of age	ask a doctor

Other information

- **each softgel contains:** sodium 6 mg

- store at controlled room temperature 15º - 30º C (59º- 86º F)
- do not use if imprinted safety seal under cap is broken or missing.

Inactive Ingredients

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

Principal Display Panel

†Compare to Dulcolax® Stool Softener active ingredient

STOOL SOFTENER

DOCUSATE SODIUM 100 mg

†This product is not manufactured or distributed by Boehringer Ingelheim Consumer Healthcare, owner of the registered trademark of Dulcolax®.

Another quality product distributed by McKesson

one post street, San Francisco, CA 94104

Money back gurantee

Please visit us at www.sunmarkbrand.com

Questions or comments? Call toll free **1-877-753-3935**

Product Label

docusate sodium 100 mg

STOOL SOFTENER

docusate sodium capsule

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49348-917
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

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 GLYCOLS (UNII: 3WJQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	

Product Characteristics			
Color	ORANGE	**Score**	no score
Shape	OVAL	**Size**	13mm
Flavor		**Imprint Code**	P51
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:49348-917-05	1 in 1 BOX		
1		25 in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part334	08/24/2010	

Labeler - Mckesson (Sunmark) (177667227)