STOOL SOFTENER LAXATIVE EXTRA STRENGTH- docusate sodium capsule, liquid filled Cardinal Health (Leader) 70000

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 (1-800-222-1222) 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^oC(68^o-77^oF); excursions permitted between 15-30^oC (59^o-86^oF)

Inactive Ingredients

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

Questions or comments?

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

Principal Display Panel

Extra Strength

Stool Softener Laxative

Docusate Sodium, 250 mg

Stool Softener Laxative

Stimulant-Free

Relieves Occasional Constipation

SOFTGELS

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

DISTRIBUTED BY CARDINAL HEALTH

DUBLIN, OHIO 43017

www.myleader.com 1-800-200-6313

Product Label



STOOL SOFTENER L	AVATI\/E EVT		.n			
docusate sodium capsule, lic		KA SI KENGI	п			
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Product Information						
Product Type	HUMAN OTC DRUG	Item Code (So	Item Code (Source) NDC		C:70000-0530	
Route of Administration	ORAL					
A ativa la avadia at/A ativa						
Active Ingredient/Active	trength					
-	Ingredient Name				_	
DOCUSATE SODIUM (UNII: F05Q)	DOCUSATE SODIUM (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG) DOCUSATE SODIU				250 mg	
Inactive Ingredients						
	Ingredient Na	ne		S	trength	
FD&C RED NO. 40 (UNII: WZ B912	27XOA)					
FD&C YELLOW NO. 6 (UNII: H77	VEI93A8)					
GELATIN (UNII: 2G86QN327L)						
GLYCERIN (UNII: PDC6A3C0OX)						
POLYETHYLENE GLYCOL, UNSP		DW1A)				
PROPYLENE GLYCOL (UNII: 6DCS	Q167V3)					
WATER (UNII: 059QF0K00R)						
SORBITOL (UNII: 506T60A25R)						
SORBITAN (UNII: 6092ICV9RU)						
MANNITOL (UNII: 30WL53L36A)						
Product Characteristics						

Sł	nape		CAPSULE	Size		20mm				
Fla	Flavor			Imprint Code		P20				
Co	Contains									
P	ackaging									
#	Item Code		Package Description		Marketing Start Date	Marketing End Date				
1			L BOTTLE, PLASTIC; Type 0: Not a ation Product		11/26/2019	12/27/2024				
Marketing Information										
	Marketing Category	Ар	plication Number or Citation	Monograph	Marketing Start Date	Marketing End Date				
OT fin	C monograph no al	ot part33	34		11/26/2019	12/27/2024				

Labeler - Cardinal Health (Leader) 70000 (063997360)

Revised: 10/2022

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