STOOL SOFTENER LAXATIVE- docusate sodium capsule, liquid filled Freds Inc

Drug Facts

Active ingredient (in each softgel)

Docusate sodium 100 mg

Purpose

Stool softener laxative

Uses

- relieves occasional constipation (irregularity)
- generally produces bowel movement in 12 to 72 hours

Warnings

Do not use

if you are presently taking mineral oil, unless told to do so by a doctor.

Ask a doctor before use if you have

- stomach pain
- nausea
- vomiting
- noticed a sudden change in bowel habits that lasts 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

• take only by mouth. Doses may be taken as a single daily dose or in divided doses.

Other information

- each softgel contains:sodium 6 mg
- store at 25°C (77°F); excursions permitted between 15-30°C (59-86°F)

Inactive ingredients

D&C red #33*, edible ink, FD&C blue #1*, FD&C red #40, FD&C yellow #6, gelatin, glycerin, polyethylene glycol, propylene glycol*, purified water, sorbitan, sorbitol, titanium dioxide

*contains one or more of these ingredients

Questions or comments?

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

Principal Display Panel

[†]Compare to the active ingredient in Colace® Regular Strength Stool Softener

Softgels

STOOL SOFTENER

Docusate Sodium

Stool softener laxative

- Gentle
- Dependable
- Stimulant-free

SOFTGELS

[†]This product is not manufactured or distributed by Purdue Products L.P., distributor of Colace® Regular Strength Stool Softener.

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

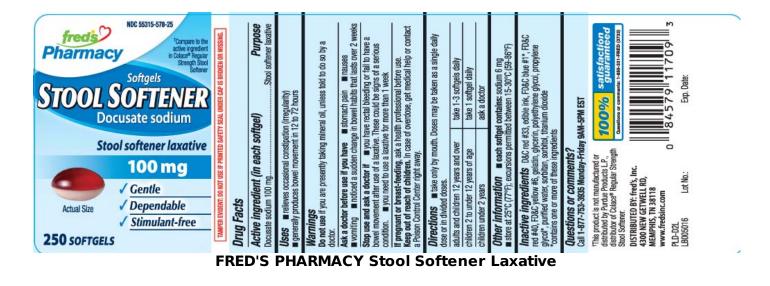
DISTRIBUTED BY: fred's, Inc.

4300 NEW GETWELL RD,

MEMPHIS, TN 38118

www.fredsinc.com

Product Label



docusate sodium capsule, liq	uid filled							
Product Information								
Product Type	HUMAN OTC DRUG	Item Code (Source) ND		NDC:553	DC:55315-578			
Route of Administration	ORAL							
Active Ingredient/Active	Moiety							
Ingr	Ingredient Name Basis of Stre				Strength			
DOCUSATE SODIUM (UNII: F05Q2	DOCUSATE SODIUM (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG)				100 mg			
Inactive Ingredients								
	S	trength						
D&C RED NO. 33 (UNII: 9DBA0SB	B0L)							
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)								
FD&C RED NO. 40 (UNII: WZ B912								
FD&C YELLOW NO. 6 (UNII: H77)	/EI93A8)							
GELATIN (UNII: 2G86QN327L)								
GLYCERIN (UNII: PDC6A3C0OX)								
POLYETHYLENE GLYCOL, UNSP	ECIFIED (UNII: 3WJQ0SDW1A	()						
PROPYLENE GLYCOL (UNII: 6DC9	Q167V3)							
WATER (UNII: 059QF0K00R)								
SORBITAN (UNII: 6092ICV9RU)								
SORBITOL (UNII: 506T60A25R)								
TITANIUM DIOXIDE (UNII: 15FIX9)								

Product Characteristics										
Color		red, white	Score		no score					
SI	Shape		OVAL	Size		12mm				
Flavor			Imprint Code		P10;SCU2;D1					
Contains										
Packaging										
#			Package Description		Marketing Date	Start	Marketing End Date			
1			n 1 BOTTLE, PLASTIC; Type 0: Not a bination Product		02/28/2019		06/30/2024			
2			50 in 1 BOTTLE, PLASTIC; Type 0: Not a ombination Product		02/28/2019		06/30/2024			
Marketing Information										
	Marketing A Category		Application Number or Monograph Citation		Marketing Start Date		Marketing End Date			
0	OTC Monograph Drug M00		07		02/28/2019		06/30/2024			

Labeler - Freds Inc (005866116)

Revised: 11/2023

Freds Inc