

DOCUSATE SODIUM- docusate sodium capsule
Forreal Pharmaceuticals, LLC

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Docusate Sodium 100 mg Softgels

Drug Facts

Active Ingredient (in each softgel)

Docusate Sodium 100 mg

Purpose

Stool softener

Uses

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

Warnings

Do not use if you are currently taking mineral oil, unless told to do so by a doctor

Ask a doctor before use if you have

- stomach pain, nausea or 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 stool softener laxative for more than 1 week

If pregnant or breastfeeding, 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 immediately 1 (800) 222-1222

Directions:

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

Adults and children 12 years and over	take 1-3 softgels daily
children 2 to under 12 years of age	take 1 softgel daily
children under 2 years of age	ask a doctor

Other information

- each capsule contains: **sodium 6 mg**
- **VERY LOW SODIUM**
- store at 25°C (77°F); excursions permitted between 15°C-30°C (59°F-86°F).
- keep tightly closed.

Inactive Ingredients: D&C Red # 33, Edible Ink, FD&C Red # 40, FD&C Yellow # 6, Gelatin, Glycerin, Polyethylene Glycol, Purified Water, Sorbitol Special, titanium dioxide

Questions or comments? (877) 367-3250

Package Label Principal Display Panel

NDC 81877-840-90

Forreal Pharmaceuticals LLC

YOUR SOLUTION TO BETTER HEALTHCARE.

Docusate

Sodium

Stool Softener

100 mg

90 Softgels

TAMPER EVIDENT: DO NOT USE IF SEAL IS BROKEN OR MISSING FROM BOTTLE.

Marketed By:

Forreal Pharmaceuticals, LLC

Royal Oak MI 48067

Questions or inquiries call

877-367-3250

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GTIN: 00381877840907
 SNO: E20D41000001
 EXP: 03/2022
 LOT: E20D41

DOCUSATE SODIUM

docusate sodium capsule

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81877-840
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
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	RED (Two-toned- white and clear red)	Score	no score
Shape	OVAL	Size	12mm
Flavor		Imprint Code	SCU2
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81877-840-90	90 in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/01/2026	

Labeler - Forreal Pharmaceuticals, LLC (118029197)

Revised: 5/2026

Forreal Pharmaceuticals, LLC