

**STOOL SOFTENER- docusate sodium capsule, liquid filled  
NuCarePharmaceuticals, Inc.**

*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.*

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**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**

**Ask a doctor before use if you**

- have stomach pain, nausea or vomiting
- have a sudden change in bowel habits that persists over a period of 2 weeks
- are presently taking mineral oil

**Stop use and ask a doctor if**

- you need to use a laxative longer than 1 week
- you have rectal bleeding or fail to have a bowel movement. These could be signs of a serious condition.

**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 and older: take 1-2 softgel daily until first bowel movement; 1 softgel daily thereafter, or as directed by doctor
- children under 12: consult a doctor
- do not exceed recommended dose

**Other information**

- **each softgel contains:** sodium 5 mg. very low sodium
- store at 15°C-25°C(59° F-77° F)
- keep tightly closed

- product from USA or Canada
- **Tamper Evident:** Do not use if imprinted seal under cap is missing or broken.

### Inactive ingredients

FD and C red 40, gelatin, glycerin, edible ink, PEG, propylene glycol, sorbitol special, water. May also contain D and C yellow 10, FD and C yellow 6 (sunset yellow), mannitol.

### Package Label

**NuCare Pharmaceuticals, Inc.**

NDC: 68071-3048-4  
**Docusate Sodium 100mg**  
**#40 Softgels**

Each softgel contains Docusate Sodium 100mg. Stool Softener Laxative FD & C Yellow #6 (sunset yellow)  
 Warnings: Ask a doctor before use if you have stomach pain, nausea or vomiting, have a sudden change in bowel habits that persists over a period of 2 weeks, are presently taking mineral oil. Stop use and ask a doctor if, you need to use a laxative longer than 1 week, you have rectal bleeding or fail to have a bowel movement. These could be signs of a serious condition. 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. Oval Clear-Red Softgel Printed: "401"

Product #: P0083040

GTIN 00368071304848  
 Serial# 00000000002  
 Exp. Date 00-00  
 LOT#: 000000

Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

WARNING: KEEP OUT OF REACH OF CHILDREN      STORE AT CONTROLLED TEMPERATURE 59-77°F.

## STOOL SOFTENER

docusate sodium capsule, liquid filled

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68071-3048(NDC:57896-401)
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)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SORBITOL (UNII: 506T60A25R)	
WATER (UNII: 059QF0K00R)	

D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)

FD&C YELLOW NO. 6 (UNII: H77VEI93A8)

MANNITOL (UNII: 3OWL53L36A)

### Product Characteristics

Color	red (reddish)	Score	no score
Shape	OVAL	Size	12mm
Flavor		Imprint Code	SCU1
Contains			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-3048-4	40 in 1 BOTTLE; Type 0: Not a Combination Product	03/14/2017	
2	NDC:68071-3048-6	60 in 1 BOTTLE; Type 0: Not a Combination Product	03/14/2017	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	01/01/2000	

**Labeler** - NuCarePharmaceuticals, Inc. (010632300)

### Establishment

Name	Address	ID/FEI	Business Operations
NuCare Pharmaceuticals, Inc.		010632300	repack(68071-3048)

Revised: 2/2021

NuCarePharmaceuticals, Inc.