

**STOOL SOFTENER- docusate sodium capsule, liquid filled  
Bryant Ranch Prepack**

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**gc 401**

**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**

- do not exceed recommended dose
- adults and children 12 years and older: take 1-3 softgels daily until first bowel movement; 1 softgel daily thereafter, or as directed by a doctor
- children under 12: consult a doctor

## Other information

- **each softgel contains:** sodium 7 mg. Very low sodium
- store at 59°-77°F (15°-25°C)
- keep tightly closed
- **Tamper Evident:** Do not use if imprinted seal under cap is missing or broken.

## Inactive ingredients

FD&C red #40, FD&C yellow #6 (sunset yellow), gelatin, glycerin, PEG, sorbitol special, water.

## HOW SUPPLIED

### Docosate Sodium 100mg Capsule

- NDC: 71335-0294-1: 30 Tablets in a BOTTLE
- NDC: 71335-0294-2: 100 Tablets in a BOTTLE
- NDC: 71335-0294-3: 60 Tablets in a BOTTLE
- NDC: 71335-0294-4: 120 Tablets in a BOTTLE
- NDC: 71335-0294-5: 90 Tablets in a BOTTLE
- NDC: 71335-0294-6: 180 Tablets in a BOTTLE
- NDC: 71335-0294-7: 10 Tablets in a BOTTLE
- NDC: 71335-0294-8: 28 Tablets in a BOTTLE
- NDC: 71335-0294-9: 56 Tablets in a BOTTLE
- NDC: 71335-0294-0: 18 Tablets in a BOTTLE

Repackaged/Relabeled by:  
Bryant Ranch Prepack, Inc.  
Burbank, CA 91504

### Docosate Sodium 100mg Capsule



GTIN 00371335029418  
Lot 208620  
Exp 7/9/2026  
SN 0123456789

Drug Facts	
<b>Active ingredient (in each softgel)</b> Docosate Sodium 100 mg	<b>Purpose</b> Stool Softener
<b>Uses</b> •Relieves occasional constipation (irregularity) •Generally produces bowel movement in 12 to 72 hours	
<b>Warnings</b> 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.	
<b>Other Information</b> • Each softgel contains: sodium 7 mg. Very low sodium • Store at 59°-77°F (15°-25°C) • Keep tightly closed • Tamper Evident: Do not use if imprinted seal under cap is missing or broken.	
<b>Directions</b> • Do not exceed recommended dose • Adults and children 12 years and older: take 1-3 softgels daily, usually 1 softgel daily after the first bowel movement, or as directed by a doctor. • Children under 12: ask a doctor	
<b>Inactive Ingredients</b> FD&C red #40, FD&C yellow #6 (sunset yellow), gelatin, glycerol, PEG, sorbitol special, water.	

NDC 71335-0294-1

Docosate Sodium

100 mg

30 Capsules



Repackaged by:  
Bryant Ranch Prepack, Inc.  
Burbank, CA 91504 USA

Manufactured by:  
Geri-Care  
Pharmaceutical Corp



7133502941

**STOOL SOFTENER**

docusate sodium capsule, liquid filled

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:71335-0294(NDC:57896-401)
<b>Route of Administration</b>	ORAL		

## Active Ingredient/Active Moiety

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>DOCUSATE SODIUM</b> (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG)	DOCUSATE SODIUM	100 mg

## Inactive Ingredients

<b>Ingredient Name</b>	<b>Strength</b>
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>GELATIN, UNSPECIFIED</b> (UNII: 2G86QN327L)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>POLYETHYLENE GLYCOL 400</b> (UNII: B697894SGQ)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>WATER</b> (UNII: 059QF0K00R)	
<b>FD&amp;C YELLOW NO. 6</b> (UNII: H77VEI93A8)	
<b>MANNITOL</b> (UNII: 3OWL53L36A)	

## Product Characteristics

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

## Packaging

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:71335-0294-1	30 in 1 BOTTLE; Type 0: Not a Combination Product	04/11/2018	
2	NDC:71335-0294-2	100 in 1 BOTTLE; Type 0: Not a Combination Product	02/27/2018	
3	NDC:71335-0294-3	60 in 1 BOTTLE; Type 0: Not a Combination Product	05/23/2018	
4	NDC:71335-0294-4	120 in 1 BOTTLE; Type 0: Not a Combination Product	02/09/2022	
5	NDC:71335-0294-5	90 in 1 BOTTLE; Type 0: Not a Combination Product	05/23/2018	
6	NDC:71335-0294-6	180 in 1 BOTTLE; Type 0: Not a Combination Product	02/09/2022	
7	NDC:71335-0294-7	10 in 1 BOTTLE; Type 0: Not a Combination Product	02/09/2022	
8	NDC:71335-0294-8	28 in 1 BOTTLE; Type 0: Not a Combination Product	09/20/2019	

9	NDC:71335-0294-9	56 in 1 BOTTLE; Type 0: Not a Combination Product	02/09/2022	
10	NDC:71335-0294-0	50 in 1 BOTTLE; Type 0: Not a Combination Product	05/18/2018	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	01/01/2000	

**Labeler** - Bryant Ranch Prepack (171714327)

**Registrant** - Bryant Ranch Prepack (171714327)

## Establishment

Name	Address	ID/FEI	Business Operations
Bryant Ranch Prepack		171714327	REPACK(71335-0294) , RELABEL(71335-0294)

Revised: 7/2024

Bryant Ranch Prepack