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

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

- 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 using a laxative. These could be a sign of a serious condition.
- you need to use a stool softener laxative for more than 1 week

If pregnant or breast-feeding,

ask a health care professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center (1-800-2222-1222) right away.

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	ask a doctor

Other information

- **each softgel contains:** sodium 7 mg
- store at 25ºC (77ºF); excursion permitted between 15-30ºC (59-86ºF)

Inactive ingredients

citric acid, FD&C Red #40, FD&C Yellow #6, gelatin, glycerin, polyethylene glycol, propylene glycol, purified water sorbitol special, white edible ink

Questions or comments?

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

HOW SUPPLIED

Docusate Sodium 100 mg

NDC: 71335-9731-1: 30 Tablets in a BOTTLE

NDC: 71335-9731-2: 100 Tablets in a BOTTLE

NDC: 71335-9731-3: 60 Tablets in a BOTTLE

NDC: 71335-9731-4: 120 Tablets in a BOTTLE

NDC: 71335-9731-5: 90 Tablets in a BOTTLE

NDC: 71335-9731-6: 180 Tablets in a BOTTLE

NDC: 71335-9731-7: 10 Tablets in a BOTTLE

NDC: 71335-9731-8: 28 Tablets in a BOTTLE

NDC: 71335-9731-9: 56 Tablets in a BOTTLE

NDC: 71335-9731-0: 18 Tablets in a BOTTLE

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

Docusate Sodium 100 mg Capsule

Each soft gelatin capsule contains:
Docusate Sodium, USP 100 mg.

Store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F).

Keep this and all drugs out of the reach of children.

Tamper evident: do not use if imprinted safety seal cap is broken or missing.

NDC 71335-**9731**-1

Docusate Sodium

100 mg

BRP PHARMACEUTICALS

30 Capsules

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

Manufactured by:
Major Pharmaceuticals

STOOL SOFTENER LAXATIVE

docusate sodium capsule, liquid filled

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71335-9731(NDC:0904-7280)
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)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SORBITAN (UNII: 6O92ICV9RU)	

Product Characteristics

Color	red	**Score**	no score
Shape	CAPSULE (Oval)	**Size**	13mm
Flavor		**Imprint Code**	PC1
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71335-9731-1	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/18/2023	
2	NDC:71335-9731-2	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/09/2023	
3	NDC:71335-9731-3	60 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/17/2023	
4	NDC:71335-9731-4	120 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2024	
5	NDC:71335-9731-5	90 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/17/2023	
6	NDC:71335-9731-6	180 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2024	
7	NDC:71335-9731-7	10 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2024	
8	NDC:71335-9731-8	28 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2024	
9	NDC:71335-9731-9	56 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2024	
10	NDC:71335-9731-0	18 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	11/15/2022	

Labeler - Bryant Ranch Prepack (171714327)

Registrant - Bryant Ranch Prepack (171714327)

Establishment

Name	Address	ID/FEI	Business Operations
Bryant Ranch Prepack		171714327	REPACK(71335-9731) , RELABEL(71335-9731)

Revised: 5/2024 Bryant Ranch Prepack