

**BISACODYL- bisacodyl tablet, delayed release
Bryant Ranch Prepack**

Major 44-327

Active ingredient (in each tablet)

Bisacodyl USP, 5 mg

Purpose

Stimulant laxative

Uses

- for relief of occasional constipation and irregularity
- this product generally produces bowel movement in 6 to 12 hours

Warnings

Do not use

if you cannot swallow without chewing.

Ask a doctor before use if you have

- stomach pain, nausea or vomiting
- a sudden change in bowel habits that lasts more than 2 weeks

When using this product

- do not chew or crush tablet(s)
- do not use within 1 hour after taking an antacid or milk
- it may cause stomach discomfort, faintness, and cramps

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

- **do not take more than directed**
- take with a glass of water

adults and children 12 years and over	take 1 to 3 tablets in a single daily dose
children 6 to under 12 years	take 1 tablet in a single daily dose
children under 6 years	ask a doctor

Other information

- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN**
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- avoid excessive humidity
- see end flap for expiration date and lot number

Inactive ingredients

acacia, ammonium hydroxide, calcium carbonate, carnauba wax, colloidal anhydrous silica, corn starch, D&C yellow #10 aluminum lake, FD&C yellow #6 aluminum lake, hypromellose, iron oxide black, lactose anhydrous, magnesium stearate, methylparaben, polydextrose, polyethylene glycol, polyvinyl acetate phthalate, povidone, propylene glycol, propylparaben, shellac glaze, simethicone, sodium alginate, sodium benzoate, sodium bicarbonate, stearic acid, sucrose, talc, titanium dioxide, triacetin, triethyl citrate

Questions or comments?

1-800-426-9391

HOW SUPPLIED

Bisacodyl USP, 5 mg

NDC: 71335-2010-0: 25 Tablets in a BOTTLE

NDC: 71335-2010-1: 30 Tablets in a BOTTLE

NDC: 71335-2010-2: 2 Tablets in a BOTTLE

NDC: 71335-2010-3: 3 Tablets in a BOTTLE

NDC: 71335-2010-4: 4 Tablets in a BOTTLE

NDC: 71335-2010-5: 10 Tablets in a BOTTLE

NDC: 71335-2010-6: 90 Tablets in a BOTTLE

NDC: 71335-2010-7: 8 Tablets in a BOTTLE

NDC: 71335-2010-8: 100 Tablets in a BOTTLE

NDC: 71335-2010-9: 20 Tablets in a BOTTLE

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

Bisacodyl 5mg Tablet

 GTIN 00371335201012 Lot 208620 Exp 3/28/2027 SN 0123456789 Package Insert	Drug Facts
	Active ingredient (in each tablet) Purpose Bisacodyl 5 mg Stimulant Laxative
	Uses •for relief of occasional constipation and irregularity. •this product generally produces bowel movement in 6 to 12 hours.
	Warnings Do not use if you cannot swallow without chewing. Ask a doctor before use if you have •stomach pain, nausea, vomiting. •noticed a sudden change in bowel habits that lasts more than 2 weeks. When using this product •do not chew or crush tablet(s). •do not use within 1 hour after taking an antacid or milk. •it may cause stomach discomfort, faintness and cramps. Stop use and ask a doctor if • you have rectal bleeding or no bowel movement after using this product. 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.
	Other Information •Tamper Evident: Do not use if imprinted seal under cap is missing or broken. •store at room temperature 15°-30°C (59°-86°F). •avoid excessive humidity •package not child resistant.
	Directions •Take with a glass of water. •Adults and children 12 years and older: take 1 to 3 tablets in a single daily dose. •Children 12 and under: ask a doctor.
Inactive Ingredients Scan Package Insert QR Code for Inactive ingredients.	

NDC 71335-2010-1

Bisacodyl Tablets, USP

5 mg

30 Tablets



Repackaged by: Bryant Ranch Prepack, Inc. **Manufactured by:** LNK International, Inc.
Burbank, CA 91504 USA



BISACODYL

bisacodyl tablet, delayed release

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71335-2010(NDC:0904-6748)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BISACODYL (UNII: 10X0709Y6I) (DEACETYLBISACODYL - UNII:R09078E41Y)	BISACODYL	5 mg

Inactive Ingredients

Ingredient Name	Strength
ACACIA (UNII: 5C5403N26O)	
AMMONIA (UNII: 5138Q19F1X)	
CALCIUM CARBONATE (UNII: H0G9379FGK)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STARCH, CORN (UNII: O8232NY3SJ)	
D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	

ANHYDROUS LACTOSE (UNII: 3S5Y5LH9PMK)
MAGNESIUM STEARATE (UNII: 70097M6I30)
METHYLPARABEN (UNII: A2I8C7HI9T)
POLYDEXTROSE (UNII: VH2XOU12IE)
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)
POLYVINYL ACETATE PHTHALATE (UNII: 58QVG85GW3)
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
PROPYLPARABEN (UNII: Z8IX2SC1OH)
SHELLAC (UNII: 46N107B71O)
DIMETHICONE (UNII: 92RU3N3Y1O)
WATER (UNII: 059QF0KO0R)
SODIUM ALGINATE (UNII: C269C4G2ZQ)
SODIUM BENZOATE (UNII: OJ245FE5EU)
SODIUM BICARBONATE (UNII: 8MDF5V39QO)
STEARIC ACID (UNII: 4ELV7Z65AP)
SUCROSE (UNII: C151H8M554)
TALC (UNII: 7SEV7J4R1U)
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)
TRIACETIN (UNII: XHX3C3X673)
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)

Product Characteristics

Color	orange	Score	no score
Shape	ROUND	Size	6mm
Flavor		Imprint Code	5
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71335-2010-0	25 in 1 BOTTLE; Type 0: Not a Combination Product	04/03/2024	
2	NDC:71335-2010-1	30 in 1 BOTTLE; Type 0: Not a Combination Product	03/03/2022	
3	NDC:71335-2010-2	2 in 1 BOTTLE; Type 0: Not a Combination Product	04/03/2024	
4	NDC:71335-2010-3	3 in 1 BOTTLE; Type 0: Not a Combination Product	04/03/2024	
5	NDC:71335-2010-4	4 in 1 BOTTLE; Type 0: Not a Combination Product	01/03/2022	
6	NDC:71335-2010-5	10 in 1 BOTTLE; Type 0: Not a Combination Product	01/18/2022	
7	NDC:71335-2010-6	90 in 1 BOTTLE; Type 0: Not a Combination Product	04/03/2024	
8	NDC:71335-2010-7	8 in 1 BOTTLE; Type 0: Not a Combination Product	01/12/2022	
9	NDC:71335-2010-8	100 in 1 BOTTLE; Type 0: Not a Combination Product	04/03/2024	
10	NDC:71335-2010-9	20 in 1 BOTTLE; Type 0: Not a Combination Product	04/03/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	12/01/2018	

Labeler - Bryant Ranch Prepack (171714327)

Registrant - Bryant Ranch Prepack (171714327)

Establishment

Name	Address	ID/FEI	Business Operations
Bryant Ranch Prepack		171714327	REPACK(71335-2010) , RELABEL(71335-2010)

Revised: 3/2025

Bryant Ranch Prepack