

**UNIT DOSE BISACODYL- bisacodyl tablet, delayed release**  
**Cardinal Health 107, LLC**

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**Major 44-327-Unit dose**

***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**
- avoid excessive humidity
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)

## ***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**

### **Distributed by:**

MAJOR® PHARMACEUTICALS

Indianapolis, IN 46268

Questions or comments?

Call (800) 616-2471

[www.majorpharmaceuticals.com](http://www.majorpharmaceuticals.com)

WARNING: This Unit Dose package is not child resistant

And is Intended for Institutional Use Only.

### **Distributed By:**

**Cardinal Health**

Dublin, OH 43017

L5071915-10325 / L5071915-20325

TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGN OF TAMPERING

Rev. 11/23	M-17	Re-order No. 301829
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## Principal Display Panel

NDC 55154-6897-0

BISACODYL USP 5 mg

Enteric Coated Tablets

10 TABLETS



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**Enteric Coated Tablets**

S105

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L5071915-10325

Lot: Exp:

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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

## UNIT DOSE BISACODYL

bisacodyl tablet, delayed release

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55154-6897(NDC:0904-6407)
Route of Administration	ORAL		

**Active Ingredient/Active Moiety**

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

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>ACACIA</b> (UNII: 5C5403N26O)	
<b>AMMONIA</b> (UNII: 5138Q19F1X)	
<b>CALCIUM CARBONATE</b> (UNII: H0G9379FGK)	
<b>CARNAUBA WAX</b> (UNII: R12CBM0EIZ)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>D&amp;C YELLOW NO. 10 ALUMINUM LAKE</b> (UNII: CQ3XH3DET6)	
<b>FD&amp;C YELLOW NO. 6 ALUMINUM LAKE</b> (UNII: GYP6Z2JR6Q)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>FERROSFERRIC OXIDE</b> (UNII: XM0M87F357)	
<b>ANHYDROUS LACTOSE</b> (UNII: 3SY5LH9PMK)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>POLYDEXTROSE</b> (UNII: VH2XOU12IE)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>POLYVINYL ACETATE PHTHALATE</b> (UNII: 58QVG85GW3)	
<b>POVIDONE, UNSPECIFIED</b> (UNII: FZ989GH94E)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>SHELLAC</b> (UNII: 46N107B71O)	
<b>DIMETHICONE, UNSPECIFIED</b> (UNII: 92RU3N3Y1O)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM ALGINATE</b> (UNII: C269C4G2ZQ)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>SODIUM BICARBONATE</b> (UNII: 8MDF5V39QO)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>SUCROSE</b> (UNII: C151H8M554)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>TRIACETIN</b> (UNII: XHX3C3X673)	
<b>TRIETHYL CITRATE</b> (UNII: 8Z96QXD6UM)	

**Product Characteristics**

<b>Color</b>	ORANGE	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	6mm
<b>Flavor</b>		<b>Imprint Code</b>	5
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55154-6897-0	10 in 1 BAG	03/25/2002	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	03/25/2002	

**Labeler** - Cardinal Health 107, LLC (118546603)

Revised: 4/2025

Cardinal Health 107, LLC