

**STIMULANT LAXATIVE- bisacodyl tablet, delayed release
BIONPHARMA INC.**

a+ health-bisacodyl DR tablets 5 mg-stimulant laxative

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

Allergy alert: Contains FD&C Yellow No. 5 (tartrazine) as a color additive.

Do not use

- if you cannot swallow without chewing

Ask a doctor before use if you have

- abdominal pain, nausea or vomiting
- noticed a sudden change in bowel habits that persists over a period of 2 weeks

Ask a doctor or pharmacist before use if you are

- taking any other drug

When using this product

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

Stop use and ask a doctor if

- you have rectal bleeding or fail to have a bowel movement after using this product. These could be signs of a serious condition.

■ you need to use for a period longer 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 use more than directed
- take with a glass of water

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

Other information

- store at 20°-25°C (68°-77°F)
- protect from excessive humidity

Inactive ingredients

colloidal silicon dioxide, FD&C yellow #5 (tartrazine), FD&C yellow #6, hypromellose, lactose monohydrate, magnesium stearate, methacrylic acid and methyl methacrylate copolymer, microcrystalline cellulose, pharmaceutical imprinting ink, sodium starch glycolate, talc, titanium dioxide, triacetin, triethyl citrate

Questions?

call **1-888-235-2466**

300's count

* **compare** to the active ingredient in **Dulcolax® Laxative**

a+ health™

stimulant laxative

**bisacodyl delayed-release
tablets USP, 5 mg**

**gentle, dependable
constipation relief**

300 tablets
enteric coated tablets

*compare to the active ingredient
in Dulcolax® Laxative

A+health™

stimulant laxative

bisacodyl delayed-release
tablets USP, 5 mg

gentle, dependable
constipation relief

300 tablets

enteric coated tablets

Do not use if printed safety seal under cap is torn or missing.

Drug Facts	
Active ingredient (in each tablet) Bisacodyl USP, 5 mg.....	Purpose Stimulant laxative
Uses	
<ul style="list-style-type: none"> ■ for relief of occasional constipation and irregularity ■ this product generally produces bowel movement in 6 to 12 hours 	
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LOT:

EXP.:

Code No.: TN/Drugs/TN00003701

PMP01840

*This product is not manufactured or distributed by Sandoz Consumer Healthcare, distributor of Dulcolax® Laxative.

Manufactured for:
BIONPHARMA
Princeton, NJ 08540
MADE IN INDIA

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PEEL HERE

T00380R0126

<p>Drug Facts (continued)</p> <p>Ask a doctor before use if you have ■ abdominal pain, nausea or vomiting ■ noticed a sudden change in bowel habits that persists over a period of 2 weeks</p> <p>Ask a doctor or pharmacist before use if you are ■ taking any other drug</p> <p>When using this product ■ it may cause stomach discomfort, faintness and cramps ■ do not chew or crush tablet(s) ■ do not use within 1 hour after taking an antacid or milk</p> <p>Stop use and ask a doctor if ■ you have rectal bleeding or fail to have a bowel movement after using this product. These could be signs of a serious condition. ■ you need to use for a period longer than 1 week</p> <p>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.</p>	<p>Directions ■ do not use more than directed</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 30%;">■ take with a glass of water</td> <td style="width: 70%;">1 to 3 tablets in a single daily dose</td> </tr> <tr> <td>adults and children 12 years of age and over</td> <td>1 tablet in a single daily dose</td> </tr> <tr> <td>children 6 to under 12 years of age</td> <td>ask a doctor</td> </tr> <tr> <td>children under 6 years of age</td> <td>ask a doctor</td> </tr> </table> <p>Other information ■ store at 20°-25°C (68°-77°F) ■ protect from excessive humidity</p>	■ take with a glass of water	1 to 3 tablets in a single daily dose	adults and children 12 years of age and over	1 tablet in a single daily dose	children 6 to under 12 years of age	ask a doctor	children under 6 years of age	ask a doctor	<p>Inactive ingredients colloidal silicon dioxide, FD&C yellow #5 (tartrazine), FD&C yellow #6, hypromellose, lactose monohydrate, magnesium stearate, methacrylic acid and methyl methacrylate copolymer, microcrystalline cellulose, pharmaceutical imprinting ink, sodium starch glycolate, talc, titanium dioxide, triacetin, triethyl citrate</p> <p>Questions? call 1-888-235-2466</p>
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STIMULANT LAXATIVE

bisacodyl tablet, delayed release

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69452-541
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
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
METHACRYLIC ACID - METHYL METHACRYLATE COPOLYMER (1:1) (UNII: 74G4R6TH13)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	

Product Characteristics

Color	orange	Score	no score
Shape	ROUND (circular)	Size	5mm
Flavor		Imprint Code	0
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69452-541-83	300 in 1 BOTTLE; Type 0: Not a Combination Product	02/23/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	02/23/2026	

Labeler - BIONPHARMA INC. (079637826)

Registrant - BIONPHARMA INC. (079637826)

Establishment

Name	Address	ID/FEI	Business Operations
Ocean Healthcare Pvt Ltd		873673519	manufacture(69452-541)