

LAXATIVE- bisacodyl tablet

Bi-Mart

Active ingredient (in each tablet)

Bisacodyl USP, 5 mg

Purpose

Laxative

Uses

- for temporary 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 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 (1-800-222-1222) right away.

Directions

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

- store at 20°-25°C (68°-77°F)

Colloidal silicon dioxide, hypromellose-5, magnesium stearate, maltodextrin, methacrylic acid, microcrystalline cellulose, neelcert FD&C yellow 10 Al Lake, neelcert FD&C yellow 6 Al Lake, PEG 400, sodium starch glycolate, talc, titanium dioxide, triethyl citrate.

call toll-free **1-844-912-4012**

Distributed by

BI-MART

Eugene, OR 97402

0 71357 00707

NDC 37835-241-01

BI-MART

*Compare to the active ingredient in Dulcolax®

ENTERIC COATED BISACODYL LAXATIVE

GENTLE AND PREDICTABLE RELIEF

100 TABLETS

Drug Facts (continued)

- 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 (1-800-222-1222) right away.

Directions ■ Take with a glass of water	adults and children 12 years of age and over	take 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)

Inactive Ingredients

Celluloid silicon dioxide, hydroxybenzoyl-, magnesium stearate, maltoedextrin, methacrylic acid, microcrystalline cellulose, neelcolor FD&C yellow 10 AI Lake, neelcolor FD&C yellow 6 AI lake, PEG 400, sodium starch glycolate, talc, titanium dioxide, triethyl citrate.

Questions? call toll-free 1-844-912-4012

*This product is not manufactured or distributed by the owner of the registered trademark Dulcolax®

DULCOLAX® (BISACODYL) AND BISMALAX® ARE REGISTERED TRADEMARKS OF THE SLOAN-PETERSON COMPANY, A DIVISION OF THE SLOAN-PETERSON GROUP, INC., 10000 WILLOW CREEK DRIVE, EUGENE, OR 97402

LAXATIVE				
bisacodyl tablet				
Product Information				
Product Type		HUMAN OTC DRUG	Item Code (Source)	NDC:37835-241
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
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				
HYPROMELLOSE 2910 (5 MPA.S) (UNII: R75537T0T4)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
MALTODEXTRIN (UNII: 7CVR7L4A2D)				
METHACRYLIC ACID (UNII: 1CS02G8656)				
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)				
D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6)				
FD&C YELLOW NO. 6 ALUMINUM LAKE (UNII: GYP6Z2JR6Q)				
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)				
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)				
TALC (UNII: 7SEV7J4R1U)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)				
Product Characteristics				
Color	orange	Score	no score	
Shape	ROUND	Size	6mm	
Flavor		Imprint Code	B5	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37835-241-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2025	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC Monograph Drug	505G(a)(3)	05/01/2025	
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Labeler - Bi-Mart (027630078)

Registrant - Bi-Mart (027630078)

Establishment			
Name	Address	ID/FEI	Business Operations
ELYSIUM PHARMACEUTICALS LIMITED		915664486	manufacture(37835-241)

Revised: 5/2025

Bi-Mart