GENTLE LAXATIVE- bisacodyl tablet, delayed release Bi-Mart

Active ingredient (in each tablet)

Bisacodyl 5 mg

Purpose

Stimulant laxative

Uses

- for relief of occasional constipation (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

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

When using this product

- do not chew or crush tablet(s)
- do not use within 1 hour after taking an antacid or milk
- you may have 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 may 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

• take with a glass of water

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

Other information

- do not use if seal around tablet is broken or damaged
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)

Inactive ingredients

anhydrous lactose, carmine, colloidal silicon dioxide, corn starch, FD&C red #40 aluminum lake, hypromellose, magnesium stearate, methacrylic acid, polydextrose, polyethylene glycol, shellac wax, simethicone, sodium bicarbonate, sodium lauryl sulfate, stearic acid, talc, titanium dioxide, triacetin, triethyl citrate.

Questions?

call toll-free 1-844-912-4012

Principal display panel



GENTLE LAXATIVE

bisacodyl tablet, delayed release

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:3783	35-244
Route of Administration	ORAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis	of Strength	Strength

Inactive Ingredients	
Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z 6XBU4)	
STARCH, CORN (UNII: 08232NY3SJ)	
FD&C RED NO. 40 ALUMINUM LAKE (UNII: 6T47AS764T)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
METHACRYLIC ACID (UNII: 1CS02G8656)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
SHELLAC (UNII: 46N107B710)	
DIMETHICONE (UNII: 92RU3N3Y10)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)	

Product Characteristics			
Color	pink	Score	no score
Shape	ROUND	Size	8mm
Flavor		Imprint Code	В
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37835- 244-30	2 in 1 CARTON	08/01/2025	
1		15 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information			
Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	08/01/2025	

Labeler - Bi-Mart (027630078)

Registrant - Bi-Mart (027630078)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(37835-244)

Revised: 8/2025 Bi-Mart