

GENTLE LAXATIVE- bisacodyl tablet

CARDINAL HEALTH

ldr441

Active ingredient (in each tablet)

Bisacodyl 5 mg

Purpose

Stimulant Laxative

Uses

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

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

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 years and under: ask a doctor

- Tamper Evident: Do not use if imprinted seal under cap is missing or broken
- store at room temperature
- avoid excessive humidity

Acacia, Ammonium Hydroxide, Calcium Carbonate, Corn Starch, D and C Yellow NO. 10 Lake, FD and C Yellow NO. 6 Lake, Hypromellose, Iron Oxide Black, Lactose, Magnesium Stearate, Methylparaben, PEG, Polydextrose, Polyvinyl Acetate Phthalate, Propylparaben, Propylene Glycol, Povidone, Shellac, Simethicone, Silica, Sodium Alginate, Sodium Benzoate, Sodium Bicarbonate, Stearic Acid, Sucrose, Talc, Titanium Dioxide, Triacetin, Triethyl Citrate, Wax

LEADER²

NDC 70000-0538-1

Gentle^{*} Laxative

Bisacodyl, 5 mg | Stimulant Laxative

^{*}Among Stimulant Laxatives

100 ENTERIC COATED TABLETS

100% Money Back Guarantee

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Inactive ingredients: Acacia, Ammonium Hydroxide, Calcium Carbonate, Corn Starch, D&C Yellow #10 Lake, FD&C Yellow #6 Lake, Hypromellose, Iron Oxide Black, Lactose, Magnesium Stearate, Methylparaben, PEG, Polydextrose, Polyvinyl Acetate Phthalate, Propylparaben, Propylene Glycol, Povidone, Shellac, Simethicone, Silica, Sodium Alginate, Sodium Benzoate, Sodium Bicarbonate, Stearic Acid, Sucrose, Talc, Titanium Dioxide, Triacetin, Triethyl Citrate, Wax.

Questions or comments? 1-800-540-3765

DIST BY CAH, DUBLIN, OH 43017 1-800-200-6313
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CIN 5579701 REV. 11/19

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bisacodyl tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70000-0538
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
AMMONIA (UNII: 5138Q19F1X)			
CALCIUM CARBONATE (UNII: H0G9379FGK)			

ACACIA (UNII: 5C5403N26O)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
STARCH, CORN (UNII: O8232NY3SJ)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
FERROSO FERRIC OXIDE (UNII: XM0M87F357)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I3O)	
METHYL PARABEN (UNII: A2I8C7HI9T)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ACETATE PHTHALATE (UNII: 58QVG85GW3)	
PROPYL PARABEN (UNII: Z8IX2SC1OH)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POVIDONE (UNII: FZ989GH94E)	
SHELLAC (UNII: 46N107B71O)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
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:70000-0538-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2019	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	11/01/2019	

Labeler - CARDINAL HEALTH (063997360)

Registrant - Geri-Care Pharmaceutical Corp (611196254)

Revised: 12/2023

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