

GENTLE LAXATIVE STIMULANT LAXATIVE- bisacodyl tablet, coated
QUALITY CHOICE (CHAIN DRUG MARKETING ASSOCIATION)

Quality Choice Gentle Laxative 25 Tablets

Active ingredient (in each tablet)

Bisacodyl, USP 5 mg

Purpose

Stimulant laxative

Use

- for temporary relief of occasional constipation and irregularity
- this product generally produces a 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 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 pregnancy 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. (1-

800-222-1222).

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	take 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

acacia, anhydrous calcium sulfate, anhydrous lactose, carnauba wax, colloidal silicon dioxide, corn starch, D&C yellow #10 aluminum lake, FD&C yellow #6 aluminum lake, gelatin, iron oxide, iron oxide black, iron oxide yellow, magnesium stearate, microcrystalline cellulose, polyethylene glycol (PEG) 400, polyvinyl acetate phthalate, povidone, shellac, sodium starch glycolate, stearic acid, sugar, talc, titanium dioxide.

Questions or comments?

1-866-467-2748

Principal Display Panel

NDC# **63868-908-25**

*Compare to the Active Ingredient in Dulcolax®

Gentle Laxative

Relieves Constipation & Irregularity

Bisacodyl, USP 5 mg

Relieves Constipation And Irregularity Overnight

Comfort-Coated Tablets For Gentle, Predictable Constipation Relief

25 TABLETS

100% QC SATISFACTION GUARANTEED

*This product is not manufactured or distributed by Boehringer Ingelheim Pharmaceuticals, Inc., the owner of the registered trademark Dulcolax®.

TAMPER EVIDENT: DO NOT USE IF any individual unit is broken or open.

Distributed by C.D.M.A., Inc.

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Novi, MI 48376-0995

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Questions: 248-449-9300

Product Label



QC(CDMA) Gentle Laxative

GENTLE LAXATIVE STIMULANT LAXATIVE

bisacodyl tablet, coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-908
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	
ACACIA (UNII: 5C5403N26O)				
CALCIUM SULFATE ANHYDROUS (UNII: E934B3V59H)				
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)				
CARNAUBA WAX (UNII: R12CBM0EIZ)				
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				
STARCH, CORN (UNII: O8232NY3S))				
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)				
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)				
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)				
FERROSOFERRIC OXIDE (UNII: XM0M87F357)				
FERRIC OXIDE RED (UNII: 1K09F3G675)				
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)				
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)				
Polyvinyl Acetate Phthalate (UNII: 58QVG85GW3)				
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)				
SHELLAC (UNII: 46N107B71O)				
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)				
STEARIC ACID (UNII: 4ELV7Z65AP)				
SUCROSE (UNII: C151H8M554)				
TALC (UNII: 7SEV7J4R1U)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				
Product Characteristics				
Color	ORANGE	Score	no score	
Shape	ROUND	Size	6mm	
Flavor		Imprint Code	RP116	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-908-25	1 in 1 CARTON	04/19/2019	
1		25 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	M007	04/19/2019	

Labeler - QUALITY CHOICE (CHAIN DRUG MARKETING ASSOCIATION) (011920774)

Revised: 11/2023

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