

WOMENS GENTLE LAXATIVE- bisacodyl tablet
TARGET CORPORATION

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

TARGET 151R 25 COUNT BLISTERS

ACTIVE INGREDIENT

In each tablet: Bisacodyl 5mg

INACTIVE INGREDIENTS

Acacia, Anhydrous Calcium Sulfate, Anhydrous Lactose, Carnauba Wax, Colloidal Silicon Dioxide, Corn Starch, D&C Red #27 Aluminum Lake, FD&C Blue #2 Aluminum Lake, FD&C Yellow #6 Aluminum Lake, Gelatin, Iron Oxide, Iron Oxide Black, Iron Oxide Yellow (Iron Oxide Ochre), Magnesium Stearate, Microcrystalline Cellulose, Polyethylene Glycol (PEG) 400, Polyvinyl Acetate Phthalate, Povidone, Shellac, Sodium Starch Glycolate, Stearic Acid, Sugar, Talc, Titanium Dioxide

Stimulant Laxative

Keep out of Reach of Children: In case of overdose, get medical help or contact a Poison Control Center right away.

Temporarily relieves occasional constipation and irregularity

Warning: 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 two weeks.

When using this product: Do not chew or crush tablet(s); Do not use within 1 hour after taking an antacid or milk; Do not use this product if you have stomach discomfort, faintness or cramps.

adults and children 12 years and over: take 1 to 3 tablets (usually 2A) daily, or as directed by a doctor

children 6 to under 12 years: take 1 tablet daily, or as directed by a doctor

children under 6 year ask a doctor

WOMENS GENTLE LAXATIVE

lot #:
exp. date:

25 ENTERIC-COATED TABLETS

upakap
women's gentle
laxative tablets
bisacodyl delayed-release
USP, 5 mg

Compare to active ingredient in Dulcolax® Pink Laxative
NDC 11025-13-1-05

ACTUAL SIZE
25
TABLETS

8160 11151

When used as directed, bisacodyl delayed-release tablets are safe and effective for the relief of occasional constipation in adults and children 12 years of age and older. The laxative effect of bisacodyl delayed-release tablets is usually experienced within 6 to 12 hours after oral administration. The laxative effect of bisacodyl delayed-release tablets is usually mild to moderate and does not cause cramping or discomfort, nor is it speeded this way.

Important Information: Do not take this medicine for more than 7 days unless your doctor tells you to. If you are taking this medicine for more than 7 days, you should see your doctor. If you are taking this medicine for more than 7 days and you do not feel better, you should see your doctor. Do not take this medicine if you are pregnant or breastfeeding. Do not take this medicine if you are taking other medicines that may interact with it. Do not take this medicine if you are taking other medicines that may interact with it. Do not take this medicine if you are taking other medicines that may interact with it.

245 05 010 NO. C00374-012
7 49483 1512
Call by Taylor Corporation
Morgantown, WV 25003
Made in U.S.A.

Drug Facts (continued) (see important information on the reverse side of this package)

Warnings: Do not take this medicine if you are pregnant or breastfeeding. Do not take this medicine if you are taking other medicines that may interact with it. Do not take this medicine if you are taking other medicines that may interact with it. Do not take this medicine if you are taking other medicines that may interact with it.

Directions: See important information on the reverse side of this package. Take 1 or 2 tablets orally with a glass of water once daily at bedtime. If you are taking this medicine for more than 7 days, you should see your doctor.

Other information: See important information on the reverse side of this package. This medicine may contain some inactive ingredients. Some of these may cause allergic reactions or other problems. Tell your doctor if you have ever had an allergic reaction to any medicine. Tell your doctor if you are taking any other medicines. Tell your doctor if you are pregnant or breastfeeding.

Drug Facts (continued) (see important information on the reverse side of this package)

Directions: See important information on the reverse side of this package. Take 1 or 2 tablets orally with a glass of water once daily at bedtime. If you are taking this medicine for more than 7 days, you should see your doctor.

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25
TABLETS

B 2 3 2 7

bisacodyl tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-151
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BISACODYL (UNII: 10 X0709Y6I) (DEACETYLBISACODYL - UNII:R09078E41Y)	BISACODYL	5 mg

Inactive Ingredients

Ingredient Name	Strength
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POVIDONE (UNII: FZ989GH94E)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
POLYVINYL ACETATE PHTHALATE (UNII: 58QVG85GW3)	
RAW SUGAR (UNII: 8M707QY5GH)	
ACACIA (UNII: 5C5403N26O)	
FERROSFERRIC OXIDE (UNII: XM0M87F357)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
STARCH, CORN (UNII: O8232NY3SJ)	
D&C RED NO. 27 (UNII: 2LRS185U6K)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GELATIN (UNII: 2G86QN327L)	
FERROUS OXIDE (UNII: G7036X8B5H)	
CALCIUM SULFATE ANHYDROUS (UNII: E934B3V59H)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SHELLAC (UNII: 46N107B71O)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	pink	Score	no score
Shape	ROUND	Size	5mm
Flavor		Imprint Code	TCL
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-151-52	25 in 1 CARTON; Type 0: Not a Combination Product	04/26/2019	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	04/01/2019	

Labeler - TARGET CORPORATION (006961700)

Registrant - TIME CAP LABORATORIES, INC. (037005209)

Establishment

Name	Address	ID/FEI	Business Operations
TIME CAP LABORATORIES, INC		037052099	manufacture(11673-151)

Revised: 4/2019

TARGET CORPORATION