BISACODYL- biscodyl tablet, coated
TARGET CORPORATION

Target Bisacodyl Tablets 5 mg

Drug Facts

Active ingredient (in each tablet)

Bisacodyl USP, 5 mg

Purpose

Stimulant Laxative

Uses

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

Warnings

WARNINGS

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.

Do not use

- if you cannot swallow without chewing

Ask a doctor before use if you have

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

When using this product

- it may cause stomach discomfort, faintness and cramps
- do not chew or crush tablet(s)
- do not use within 1 hour after taking an antacid or milk

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

Other information

- **each tablet contains:** magnesium 5 mg
- store at 25°C (77°F) excursions permitted between 15°-30°C (59°-86°F)
- use by expiration date on package
- protect from excessive humidity

Directions

take with a glass of water
- adults and children 12 years of age and over: 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

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 (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

Questions or comments?

Call **1-800-910-6874**

Drug Facts (continued)

Stop use and ask a doctor if
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Questions or comments?

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Adhesive Area

Compare to active ingredient in Dulcolax® Laxative*

Extra Gentle Laxative Tablets

Bisacodyl Delayed-Release USP, 5 mg

- Stimulant laxative
- Gentle, dependable constipation relief

up&up™

100 ENTERIC-COATED TABLETS

Actual Size

TAMPER EVIDENT: DO NOT USE THIS PRODUCT IF THE IMPRINTED FOIL SEAL OVER THE MOUTH OF THE BOTTLE IS CUT, TORN, BROKEN OR MISSING

Drug Facts

Active ingredient (in each tablet)	Purpose
Bisacodyl USP, 5 mg	Stimulant laxative

Uses

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- this product generally produces bowel movement in 6-12 hours

Warnings

Do not use
- if you cannot swallow without chewing

Ask a doctor before use if you have
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- noticed a sudden change in bowel habits that lasts more than 2 weeks

When using this product
- it may cause stomach discomfort, faintness and cramps
- do not chew or crush tablet(s)
- do not use within 1 hour after taking an antacid or milk

Stop use and ask a doctor if
- you have rectal bleeding or fail to have a bowel

Drug Facts (continued under label)

*This product is not manufactured or distributed by Sanofi Consumer Healthcare, distributor of Dulcolax® Laxative.

NDC 11673-182-01
245 05 55549 R01
C-002262-01-073-0000
Dist. by Target Corp.
Minneapolis, MN 55403
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Target Brands, Inc.
14R 0624

3 11673 18201 7

PEEL HERE

LOT:

EXP.:

Varnish Omit Area

Drug Facts (continued)

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 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

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TAMPER EVIDENT: EACH TABLET IS SAFETY SEALED. DO NOT USE IF BLISTER OR FOIL IS BROKEN OR MISSING

LOT #:

EXP. DATE:

Varnish Omit Area

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Manufactured or distributed by Sanofi Consumer Healthcare, distributor of Dulcolax® Laxative

Questions or comments? Call 1-800-910-6874

NDC C-002262-01-073-0000

Dist by Target Corporation

Minneapolis, MN 55403

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14R 0324

BISACODYL

biscodyl tablet, coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-182
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: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
POVIDONE (UNII: FZ989GH94E)	
GELATIN (UNII: 2G86QN327L)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POLYVINYL ACETATE PHTHALATE (UNII: 58QVG85GW3)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
SUCROSE (UNII: C151H8M554)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SHELLAC (UNII: 46N107B71O)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	

Product Characteristics

Color	orange	**Score**	no score
Shape	ROUND	**Size**	5mm
Flavor		**Imprint Code**	TCL003
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-182-52	25 in 1 BLISTER PACK; Type 0: Not a Combination Product	02/04/2020	
2	NDC:11673-182-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	06/15/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	02/04/2020	

Labeler - TARGET CORPORATION (006961700)

Registrant - TIME CAP LABORATORIES, INC (037052099)

Establishment

Name	Address	ID/FEI	Business Operations
TIME CAP LABORATORIES, INC.		037052099	manufacture(11673-182)

Revised: 7/2024 TARGET CORPORATION