

**GENTLE LAXATIVE- bisacodyl tablet, delayed release
BETTER LIVING BRANDS LLC.**

14R_Albertsons_21130-410_Bisacodyl Delayed Release 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

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).

Directions

- take with a glass of water

adults and children 12 years of age and over
children 6 to under 12 years of age
children under 6 years of age

1 to 3 tablets in a single daily dose
1 tablet in a single daily dose
ask a doctor

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

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? 1-877-290-4008

Bisacodyl tablets 25ct IFC

Drug Facts (continued)

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.

Drug Facts (continued)

titanium dioxide

Questions or comments? 1-877-290-4008

The product is not manufactured or distributed by Sunovion. It is manufactured by Sunovion Laboratories, Inc. (Sunovion) and is distributed by Sunovion Laboratories, Inc. (Sunovion).

Drug Facts

Active ingredient
Bisacodyl USP, 5 mg

Other ingredients
Simulant lactose

Uses
For relief of occasional constipation and irregularity in the product generally produce bowel movement in 6-12 hours

Warnings
Do not use if you cannot swallow without chewing
Ask a doctor before use if you have stomach pain, nausea or vomiting
If noticed a sudden change in bowel habits that lasts more than 2 weeks
When using this product it may cause stomach discomfort, flatness and cramps
Do not use 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.

Directions
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

Other information
Each tablet contains: magnesium 5 mg
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Drug Facts (continued)

signs of a serious condition.
If you need to use a laxative for more than 1 week, ask a health professional before use.
Keep out of reach of children. In case of overdose, get medical help or contact Poison Control Center right away (1-800-222-1222).

Directions
Adults and children 12 years of age and over: 1 to 3 tablets in a single daily dose
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SmartLabel
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P.O. BOX 99
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14R 0425




Signature SELECT

COMPARE TO
Dulcolax® Laxative Tablets
active ingredient*

NDC 21130-410-52

LOT:
EXP:

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

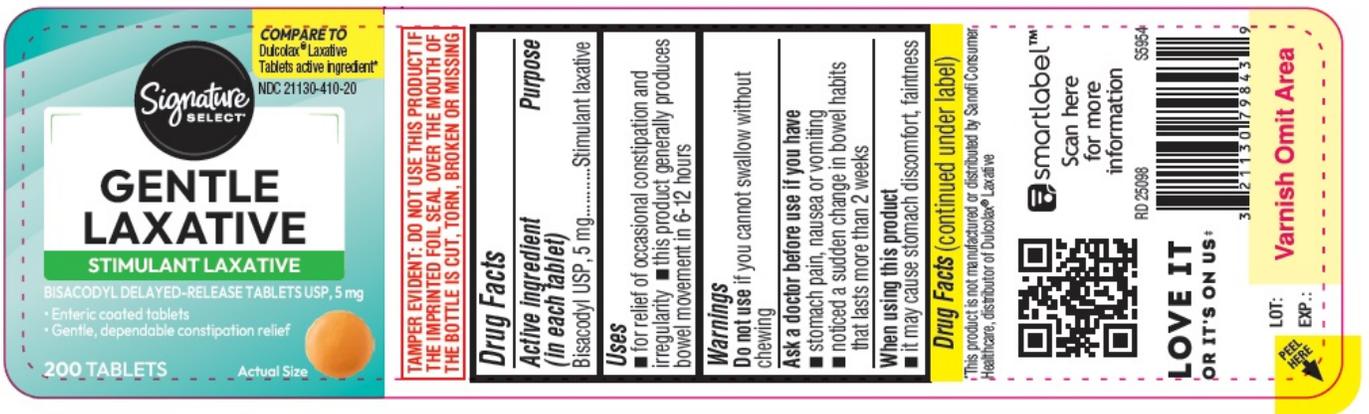
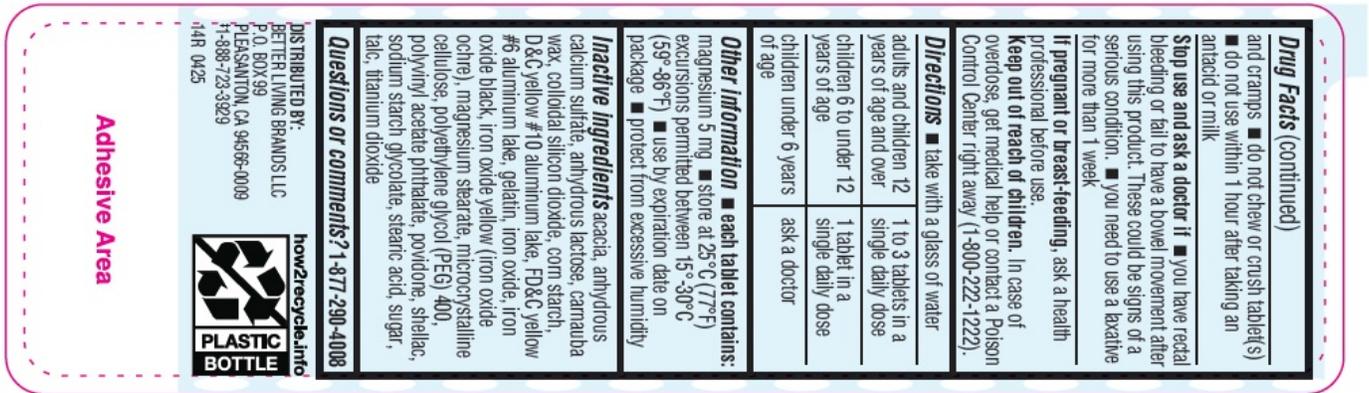
READ AND KEEP CARTON FOR COMPLETE WARNINGS AND INFORMATION

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Bisacodyl tablets 200ct label



GENTLE LAXATIVE

bisacodyl tablet, delayed release

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:21130-410

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
FERRIC OXIDE RED (UNII: 1K09F3G675)	
CALCIUM SULFATE ANHYDROUS (UNII: E934B3V59H)	
D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6)	
FERROSFERRIC OXIDE (UNII: XM0M87F357)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POLYVINYL ACETATE PHTHALATE (UNII: 58QVG85GW3)	
SHELLAC (UNII: 46N107B710)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
FD&C YELLOW NO. 6 ALUMINUM LAKE (UNII: GYP6Z2JR6Q)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SUCROSE (UNII: C151H8M554)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
ACACIA (UNII: 5C5403N26O)	
GELATIN (UNII: 2G86QN327L)	
POVIDONE (UNII: FZ989GH94E)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	orange	Score	no score
Shape	ROUND	Size	6mm
Flavor		Imprint Code	TCL003
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21130-410-52	1 in 1 CARTON	06/01/2023	
1		25 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:21130-	200 in 1 BOTTLE; Type 0: Not a Combination	06/01/2023	

410-20

Product

00/22/2023

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	06/01/2023	

Labeler - BETTER LIVING BRANDS LLC. (009137209)

Registrant - Time Cap Labs Inc. (037052099)

Establishment

Name	Address	ID/FEI	Business Operations
TIME CAP LABORATORIES, INC.		037052099	manufacture(21130-410)

Revised: 9/2025

BETTER LIVING BRANDS LLC.