

LAXATIVE- bisacodyl tablet, delayed release
Publix Super Markets Inc

Publix 44-327

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

Stop use and ask a doctor if

- you need to use a laxative for more than 1 week
- you have rectal bleeding or fail to have a bowel movement after use of a laxative.
These could be signs of a serious condition.

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.

Directions

- take with a glass of water

adults and children 12 years and over	take 1 to 3 tablets in a single daily dose
children 6 to under 12 years	take 1 tablet in a single daily dose
children under 6 years	ask a doctor

Other information

- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN**
- avoid excessive humidity
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- see end flap for expiration date and lot number

Inactive ingredients

acacia, ammonium hydroxide, calcium carbonate, carnauba wax, colloidal anhydrous silica, corn starch, D&C yellow #10 aluminum lake, FD&C yellow #6 aluminum lake, hypromellose, iron oxide black, lactose anhydrous, magnesium stearate, methylparaben, polydextrose, polyethylene glycol, polyvinyl acetate phthalate, povidone, propylene glycol, propylparaben, shellac glaze, simethicone, sodium alginate, sodium benzoate, sodium bicarbonate, stearic acid, sucrose, talc, titanium dioxide, triacetin, triethyl citrate

Principal display panel

NDC 56062-327-56

laxative

BISACODYL USP 5, mg

STIMULANT LAXATIVE

- Overnight relief
- Gentle, dependable, constipation relief

ACTUAL SIZE

25 COMFORT

COATED TABLETS

†Compare to the active ingredient in Dulcolax[®] Laxative Tablets

†This product is not manufactured or distributed by Sanofi-Aventis Deutschland GMBH, owner of the registered trademark Dulcolax[®] Laxative Tablets.

50844 ORG011932756

TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING

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PUBLIX GUARANTEE: COMPLETE SATISFACTION OR YOUR MONEY BACK



laxative

BISACODYL USP, 5 mg

STIMULANT LAXATIVE



NDC 56062-327-56

laxative

BISACODYL USP, 5 mg

STIMULANT LAXATIVE

E1029R

- Overnight relief
- Gentle, dependable, constipation relief



ACTUAL SIZE

25 COMFORT
COATED TABLETS

¹Compare to the active ingredient in Dulcolax® Laxative Tablets

Drug Facts (continued)
black, lactose anhydrous, magnesium stearate, methylparaben, polydextrose, polyethylene glycol, polyvinyl acetate phthalate, povidone, propylene glycol, propylparaben, shellac glaze, simethicone, sodium alginate, sodium benzoate, sodium

Drug Facts (continued)
bicarbonate, stearic acid, sucrose, talc, titanium dioxide, tracebn, triethyl citrate

B-1518-327-56-R
OR0011932756

0-41415-51773-8

DISTRIBUITE
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TAMPER
EVIDENT
UNIT

Other information
 ■ TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN ■ avoid excessive handling ■ store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F) ■ see end flap for expiration date and lot number

Inactive ingredients
aluminum lake, hypromellose, iron oxide yellow #10 aluminum lake, FD&C yellow #6 coiloidal anhydrous silica, corn starch, D&C hydroxide, calcium carbonate, carnauba wax, acacia, ammonium

12 years
children under 6 ask a doctor
daily dose

Do not use if you cannot swallow without chewing.

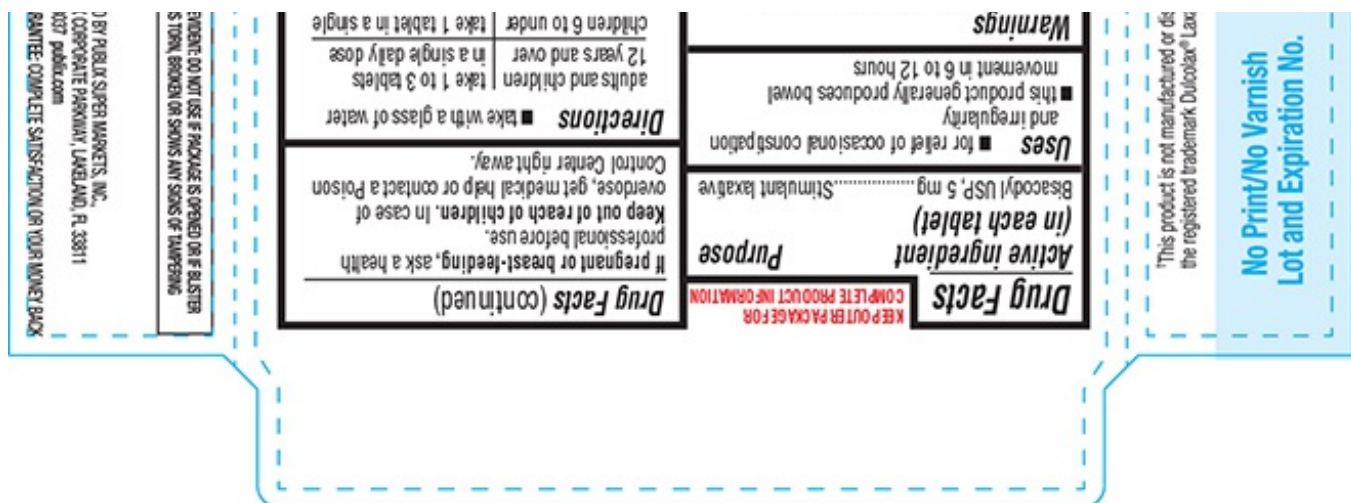
Ask a doctor before use if you have
 ■ stomach pain, nausea or vomiting
 ■ sudden change in bowel habits that lasts more than 2 weeks

When using this product
 ■ do not use within 1 hour after taking an antacid or milk ■ do not chew or crush tablet(s) ■ you may have stomach discomfort, flatness and cramps

Stop use and ask a doctor if
 ■ you need to use a laxative for more than 1 week
 ■ you have rectal bleeding or fail to have a bowel movement after use of a laxative. These could be signs of a serious condition.

distributed by Sanofi-Aventis Deutschland GmbH, owner of
active tablets. 50844 OR0011932756

Publix



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LAXATIVE

bisacodyl tablet, delayed release

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:56062-327
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)	
AMMONIA (UNII: 5138Q19F1X)	
CALCIUM CARBONATE (UNII: H0G9379FGK)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STARCH, CORN (UNII: O8232NY3SJ)	
D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
FERROSFERRIC OXIDE (UNII: XM0M87F357)	
ANHYDROUS LACTOSE (UNII: 3S5Y5LH9PMK)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
Polyvinyl Acetate Phthalate (UNII: 58QVG85GW3)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

PROPYLPARABEN (UNII: Z8IX2SC1OH)
SHELLAC (UNII: 46N107B71O)
DIMETHICONE (UNII: 92RU3N3Y1O)
WATER (UNII: 059QF0KO0R)
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:56062-327-56	1 in 1 CARTON	09/15/2021	
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	505G(a)(3)	09/15/2021	

Labeler - Publix Super Markets Inc (006922009)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	manufacture(56062-327) , pack(56062-327)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(56062-327) , pack(56062-327)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(56062-327)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(56062-327)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(56062-327)

Revised: 10/2023

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