RIGHT REMEDIES OVERNIGHT LAXATIVE- bisacodyl tablet Strive Pharmaceuticals Inc.

RIGHT REMEDIES OVERNIGHT Laxative BISACODYL (USP) 5 mg

Drug Facts

Active ingredient (in each tablet)

Bisacodyl (USP) 5 mg

Purpose

Stimulant laxative

Use

- 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
- 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.

Directions

- do not take more than directed
- take with a glass of water

adults and	1 to 3 tablets in a single daily
children	dose. It is recommended to
12 years	start with the lowest dose (1
of age and	tablet) and increasing the next
over	day as needed
children 6	1 tablet in a single daily dose
to under	
12 years	
of age	
children	ask a doctor
under 6	
years	

Other information

- contains FD&C Yellow No. 6
- store at room temperature between 20°-25°C (68°-77°F)
- protect from light, heat and moisture

Inactive ingredients

acacia gum, calcium carbonate, croscarmellose sodium, D&C Yellow No.10 aluminum lake, dibasic calcium phosphate, FD&C Yellow No. 6 aluminum lake, gelatin, glycerol monostearate, hypromellose, iron oxide red, magnesium stearate, methacrylic acid copolymer type C, microcrystalline cellulose, polyethylene glycol, precipitated silica, silicon dioxide, sodium methylparaben, sodium propylparaben, sodium starch glycolate, starch, sucrose, talcum, titanium dioxide, triethyl citrate

Questions or comments?

1-888-577-8033 Monday - Friday 8am - 4pm EST

Compare to the active ingredient in **Dulcolax**® Laxative Tablets*

Gentle, effective overnight relief of occasional constipation

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

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

Distributed by: Strive Pharmaceuticals Inc., East Brunswick, NJ 08816

PRODUCT OF INDIA

Packaging



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Drug Facts Label

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Drug Facts (continued)

- 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.

Directions

- do not take more than directed
- take with a glass of water

adults and children 12 years of age and over	1 to 3 tablets in a single daily dose. It is recommended to start with the lowest dose (1 tablet) and increasing the next day as needed
children 6 to under 12 years of age	1 tablet in a single daily dose
children under 6 years	ask a doctor

Other information

- contains FD&C Yellow No. 6
- store at room temperature between 20°-25°C (68°-77°F)
- protect from light, heat and moisture

Inactive ingredients
acacia gum, calcium carbonate, croscarmellose sodium,
D&C Yellow No.10 aluminum lake, dibasic calcium phosphate, FD&C Yellow No. 6 aluminum lake, gelatin, glycerol monostearate, hypromellose, iron oxide red, magnesium stearate, methacrylic acid co-polymer type C, microcrystalline cellulose, polyethylene glycol, precipitated silica, silicon dioxide, sodium methylparaben, sodium propylparaben, sodium starch glycolate, starch, sucrose, talcum, titanium dioxide, triethyl citrate

Questions or comments? 1-888-577-8033 Monday - Friday 8am - 4pm EST

RIGHT REMEDIES OVERNIGHT LAXATIVE

bisacodyl tablet

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70692-761	
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 CARBONATE (UNII: H0G9379FGK)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6)	
ANHYDROUS DIBASIC CALCIUM PHOSPHATE (UNII: L11K75P92J)	
FD&C YELLOW NO. 6 ALUMINUM LAKE (UNII: GYP6Z2JR6Q)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
METHACRYLIC ACID AND ETHYL ACRYLATE COPOLYMER (UNII: NX76LV5T8J)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
METHYLPARABEN SODIUM (UNII: CR6K9C2NHK)	
PROPYLPARABEN SODIUM (UNII: 625NNB0G9N)	

SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)	
STARCH, CORN (UNII: O8232NY3SJ)	
SUCROSE (UNII: C151H8M554)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)	

Product Characteristics				
Color	orange	Score	no score	
Shape	ROUND	Size	6mm	
Flavor		Imprint Code	G	
Contains				

	Packaging					
# Item Code Package Description		le Package Description	Marketing Start Date	Marketing End Date		
	1 NDC:70692-7	61- 300 in 1 BOTTLE; Type 0: Not a Combination Product	04/25/2025			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	505G(a)(3)	04/25/2025		

Labeler - Strive Pharmaceuticals Inc. (080028013)

Revised: 4/2025 Strive Pharmaceuticals Inc.