

**DULCOLAX STIMULANT LAXATIVE- bisacodyl tablet, coated**

**Chattem, Inc.**

**Reference Label Set Id: e0dcae0f-6dca-4700-a415-de2274e6f779**

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**Dulcolax Stimulant Laxative**

***Dulcolax Stimulant Laxative Tablets***

***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**

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take with a glass of water

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adults and children 12 years of age and over 1 to 3 tablets in a single daily dose

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children 6 to under 12 years of age 1 tablet in a single daily dose

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children under 6 years of age ask a doctor

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**Other information**

- contains FD&C Yellow No. 6
- do not use if individual blister unit is open or torn
- store at 20°-25°C (68°-77°F)
- protect from excessive humidity

**Inactive ingredients**

acacia senegal gum, 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 oxides, lactose, magnesium stearate, methylparaben, polydextrose, polyethylene glycol, polyvinyl acetate phthalate, povidone, propylene glycol, propylparaben, shellac, simethicone, sodium alginate, sodium benzoate, sodium bicarbonate, stearic acid, sucrose, talc, titanium dioxide, triacetin, triethyl citrate

**Questions?**

Call **1-866-844-2798** or visit **www.Dulcolax.com**

**Keep carton as it contains important product information.**

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**PRINCIPAL DISPLAY PANEL**

Dulcolax  
LAXATIVE  
200 TABLETS





## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:41167-0400
<b>Route of Administration</b>	ORAL		

## Active Ingredient/Active Moiety

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

## Inactive Ingredients

<b>Ingredient Name</b>	<b>Strength</b>
<b>ACACIA</b> (UNII: 5C5403N26O)	
<b>CALCIUM CARBONATE</b> (UNII: H0G9379FGK)	
<b>CARNAUBA WAX</b> (UNII: R12CBM0EIZ)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>FD&amp;C YELLOW NO. 6</b> (UNII: H77VEI93A8)	
<b>D&amp;C YELLOW NO. 10</b> (UNII: 35SW5USQ3G)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)	
<b>LACTOSE, UNSPECIFIED FORM</b> (UNII: J2B2A4N98G)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>POLYDEXTROSE</b> (UNII: VH2XOU12IE)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>Polyvinyl Acetate Phthalate</b> (UNII: 58QVG85GW3)	
<b>POVIDONE</b> (UNII: FZ989GH94E)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>SHELLAC</b> (UNII: 46N107B71O)	
<b>SODIUM ALGINATE</b> (UNII: C269C4G2ZQ)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>SODIUM BICARBONATE</b> (UNII: 8MDF5V39QO)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>SUCROSE</b> (UNII: C151H8M554)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>TRIACETIN</b> (UNII: XHX3C3X673)	
<b>TRIETHYL CITRATE</b> (UNII: 8Z96QXD6UM)	

## Product Characteristics

<b>Color</b>	orange	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	5mm
<b>Flavor</b>		<b>Imprint Code</b>	DL
<b>Contains</b>			

<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41167-0400-5	1 in 1 CARTON	04/14/2019	
1		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:41167-0400-6	8 in 1 CARTON	04/14/2019	
2		25 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:41167-0400-8	3 in 1 CARTON	05/01/2019	
3		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
4	NDC:41167-0400-3	2 in 1 CARTON	04/14/2019	
4		25 in 1 BLISTER PACK; Type 0: Not a Combination Product		
5	NDC:41167-0400-2	1 in 1 CARTON	04/14/2019	
5		25 in 1 BLISTER PACK; Type 0: Not a Combination Product		
6	NDC:41167-0400-1	1 in 1 CARTON	04/14/2019	
6		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
7	NDC:41167-0400-4	4 in 1 CARTON	06/03/2019	
7		25 in 1 BLISTER PACK; Type 0: Not a Combination Product		
8	NDC:41167-0400-9	1 in 1 CARTON	06/01/2022	
8		6 in 1 BLISTER PACK; Type 0: Not a Combination Product		

<b>Marketing Information</b>			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	04/14/2019	

<b>DULCOLAX STIMULANT LAXATIVE</b>			
bisacodyl tablet, coated			
<b>Product Information</b>			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:41167-0200
<b>Route of Administration</b>	ORAL		
<b>Active Ingredient/Active Moiety</b>			
	<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>

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<b>Flavor</b>		<b>Imprint Code</b>	DL
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41167-0200-5	1 in 1 CARTON	02/25/2019	
1		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:41167-0200-6	8 in 1 CARTON	04/11/2019	

2		25 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:41167-0200-8	3 in 1 CARTON	01/02/2019	
3		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
4	NDC:41167-0200-3	2 in 1 CARTON	01/02/2019	
4		25 in 1 BLISTER PACK; Type 0: Not a Combination Product		
5	NDC:41167-0200-2	1 in 1 CARTON	02/15/2019	
5		25 in 1 BLISTER PACK; Type 0: Not a Combination Product		
6	NDC:41167-0200-1	1 in 1 CARTON	03/01/2019	
6		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
7	NDC:41167-0200-4	4 in 1 CARTON	01/02/2019	
7		25 in 1 BLISTER PACK; Type 0: Not a Combination Product		

## Marketing Information

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OTC Monograph Drug	505G(a)(3)	01/02/2019	

**Labeler** - Chattem, Inc. (003336013)

Revised: 4/2025

Chattem, Inc.