DULCOLAX STOOL SOFTENER- docusate sodium capsule, liquid filled Chattem, Inc.

Dulcolax Stool Softener - Patheon

Dulcolax® Stool Softener Carton Drug Facts

Active ingredient (in each capsule)

Docusate sodium (USP) 100 mg

Purpose

Stool softener laxative

Use

- for relief of occasional constipation and irregularity
- this product generally produces bowel movement in 12 to 72 hours

Warnings

Do not use

• if you are presently taking mineral oil, unless told to do so by a doctor

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

Stop use and ask a doctor if

● you have rectal bleeding or no 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

take with a glass of water1 to 3 capsules daily. This doseadults and children 12 years of age and over
may be taken as a single daily dose or in divided doses.1 to 3 capsules daily. This dosechildren 2 to under 12 years of age1 capsule dailychildren under 2 years of ageask a doctor

Other information

- each capsule contains: **sodium 6 mg**
- store at 20-25°C (68-77°F). Protect from excessive humidity
- contains FD&C Yellow No. 6
- do not use this product if the safety seal under the cap is torn or missing

Inactive ingredients

FD&C red no. 40, FD&C yellow no. 6, gelatin, glycerin, mannitol, pharmaceutical ink, polyethylene glycol 400, propylene glycol, sorbitan, sorbitol, water

Questions?

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

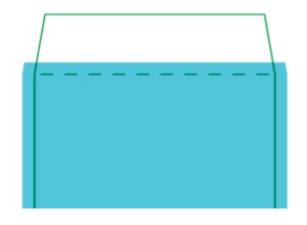
Keep carton as it contains important product information.

PRINCIPAL DISPLAY PANEL

Dulcolax

Stool Softener

25 Liquid Gels







DULCOLAX STOOL SOFTENER docusate sodium capsule, liquid filled **Product Information** HUMAN OTC DRUG NDC:41167-0221 **Product Type** Item Code (Source) **Route of Administration** ORAL **Active Ingredient/Active Moiety Ingredient Name** Basis of Strength Strength DOCUSATE SODIUM (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG) DOCUSATE SODIUM 100 mg **Inactive Ingredients Ingredient Name** Strength FD&C RED NO. 40 (UNII: WZ B9127XOA) FD&C YELLOW NO. 6 (UNII: H77VEI93A8) GELATIN (UNII: 2G86QN327L) GLYCERIN (UNII: PDC6A3C0OX) MANNITOL (UNII: 30WL53L36A) POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ) **PROPYLENE GLYCOL** (UNII: 6DC9Q167V3) SORBITAN (UNII: 6092ICV9RU) SORBITOL (UNII: 506T60A25R) WATER (UNII: 059QF0KO0R) **Product Characteristics** Color red Score no score OVAL Size 10mm Shape Flavor Dulcolax **Imprint Code** Contains

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41167- 0221-4	1 in 1 CARTON	08/01/2020	
L		100 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:41167- 0221-2	1 in 1 CARTON	08/01/2020	
2		25 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:41167- 0221-3	1 in 1 CARTON	08/01/2020	
3		50 in 1 BOTTLE; Type 0: Not a Combination Product		
1	NDC:41167- 0221-9	1 in 1 CARTON	08/01/2020	09/18/2024
l		30 in 1 BOTTLE; Type 0: Not a Combination Product		
M	larketing l	nformation		
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Dru		M007	08/01/2020	

Labeler - Chattem, Inc. (003336013)

Revised: 2/2024

Chattem, Inc.