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

Dulcolax Pink Stool Softener

Dulcolax Pink®

Stool Softener

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 water

adults and children 12 years of age and over	1 to 3 capsules daily. This dose may be taken as a single daily dose or in divided doses.
children 2 to under 12 years of age	1 capsule daily
children under 2 years of age	ask a doctor

Other information

- **each capsule contains:** sodium 6 mg
- store at 20°-25°C (68°-77°F)
- protect from excessive humidity
- do not use this product if the safety seal under the cap is torn or missing

Inactive ingredients

D&C red no. 33, FD&C blue no. 1, FD&C red no. 40, FD&C yellow no. 6, gelatin, glycerin, mannitol, pharmaceutical ink, polyethylene glycol 400, propylene glycol, sorbitan, sorbitol, titanium dioxide, water

Questions?

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

PRINCIPAL DISPLAY PANEL

Dulcolax
Pink
STOOL SOFTENER
25 SOFTGELS

DULCOLAX PINK STOOL SOFTENER

docusate sodium capsule, liquid filled

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41167-0281
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**
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
MANNITOL (UNII: 3OWL53L36A)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SORBITAN (UNII: 6O92ICV9RU)	
SORBITOL (UNII: 506T60A25R)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics			
Color	red, white	**Score**	no score
Shape	OVAL	**Size**	13mm
Flavor		**Imprint Code**	L486
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:41167-0281-1	1 in 1 CARTON	02/18/2018	
1		25 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:41167-0281-2	1 in 1 CARTON	09/07/2018	
2		30 in 1 BOTTLE; 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)	02/18/2018	

Labeler - Chattem, Inc. (003336013)

Revised: 11/2023 Chattem, Inc.