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 taken as a single daily dose or in divided doses.

1 to 3 capsules daily. This dose may be

children 2 to under 12 years of age

children under 2 years of age

1 capsule daily 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

Route of Administration ORAL Active Ingredient/Active Moiety Ingredient Name Basis of Strength Strength		Item Code (Source) NDC:41167-0281 NDC:41167-0281 NDC:41167-0281						7 0201
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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.