PHILLIPS STOOL SOFTENER BULK DOCUSATE- docusate sodium capsule, liquid filled
Bayer HealthCare LLC.

Phillips ®' Stool Softener Liquid Gels

Drug Facts

Active ingredient (in each softgel)

Docusate sodium 100 mg

Purpose

Stool softener laxative

Use

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

Warnings

Do not use if you have ever had an allergic reaction to this product or any of its ingredients.

Ask a doctor before use if you have

- stomach pain, nausea, or vomiting
- a sudden change in bowel habits that lasts over 14 days

Ask a doctor or pharmacist before use if you are presently taking mineral oil

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 softgels with a full glass (8 oz) of water

adults and children 12 years and older	take 1 to 3 softgels daily or as directed by a doctor. This dose may be taken as a

years and older	single daily dose or in divided doses.
children 6 to under 12 years	take 1 softgel daily or as directed by a doctor
children under 6 years of age	ask a doctor

Other information

- **each softgel contains:** sodium 6 mg
- very low sodium
- store at room temperature. Avoid excessive heat 40°C (104°F).

Inactive ingredients

FD&C blue #1, FD&C red #40, FD&C yellow #6, gelatin, glycerin, polyethylene glycol, povidone, propylene glycol, purified water, shellac, sodium hydroxide, sorbitol sorbitan solution, titanium dioxide

Questions or comments?

1-800-986-0369 (Mon – Fri 9AM – 5PM EST) or www.bayercare.com

PRINCIPAL DISPLAY PANEL - 30 Capsule Carton

PHILLIPS STOOL SOFTENER BULK DOCUSATE

docusate sodium capsule, liquid filled

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0280-1012
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**
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SHELLAC (UNII: 46N107B71O)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

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)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
SORBITOL (UNII: 506T60A25R)	
SORBITAN (UNII: 6O92ICV9RU)	

Product Characteristics

Color	red	**Score**	no score
Shape	OVAL	**Size**	4mm
Flavor		**Imprint Code**	P10
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0280-1012-30	1 in 1 CARTON	05/19/2015	
1		30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:0280-1012-60	1 in 1 CARTON	05/19/2015	12/01/2023
2		60 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	09/02/2013	

Labeler - Bayer HealthCare LLC. (112117283)

Revised: 10/2024 Bayer HealthCare LLC.