SILACE - docusate sodium syrup Physicians Total Care, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Silace syrup

Active Ingredient: Docusate sodium 60 mg (in each 15 mL (1 tablespoonful))

Purpose: Stool Softener

Uses

- for gentle, reliable relief from occasional constipation (irregularity)
- generally produces bowel movement in 12 to 72 hours

Warnings

- Do not use
- laxative products for longer than 1 week unless told to do so by a doctor
- **Do not use** if you are presently taking mineral oil unless told to do by a doctor

Ask a doctor before use if you have

- stomach pain
- nausea
- vomiting
- noticed a sudden change in bowel habits that last over two weeks

Stop use and ask a doctor if you have rectal bleeding or fail to have a bowel movement after use of a laxative. These could be signs of a serious condition

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 as directed by a doctor
- doses must be given in a 6-8 oz glass of milk or fruit juice, to prevent throat irritation
- dose may be taken as a single daily dose or in divided doses
- dosage should be adjusted to individual response

Adults and children 12 years of age and older	1 to 6 tablespoonfuls, or as directed by a doctor
Children 6 to under 12 years of age	1 to 2 1/2 tablespoonfuls, or as directed by a doctor
Children under 6 years	Ask a doctor

Other information

- store at room temperature 20°-25°C (68°-77°F)
- protect from freezing and excessive heat
- do not use if tamper-evident safety seal around cap is broken or missing
- dispense in tight, light-resistant container with a child-resistant closure

Inactive ingredients:

alcohol not more than 1%, citric acid, D&C red no. 33, FD&C red no. 40, peppermint flavor, glycerin, methylparaben, propylene glycol, propylparaben, purified water, sodium benzoate, sodium citrate, sucrose.

Questions

888-974-5279

Manufactured by:

Silarx Pharmaceutical, Inc.

19 West Street

Spring Valley, NY 10977-USA.

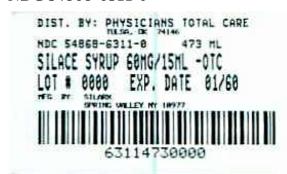
Relabeling of additional barcode by:

Physicians Total Care, Inc. Tulsa, OK 74146

SILACE (docusate sodium) syrup

473 mL (1 Pint)

NDC 54868-6311-0



SILACE

docusate sodium syrup

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:54868-6311(NDC:54838-107) Route of Administration ORAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Docusate sodium (UNII: F05Q2T2JA0) (Docusate - UNII:M7P27195AG)	Docusate sodium	60 mg in 15 mL	

Ingredient NameStrengthalcohol (UNII: 3K9958 V90M)	Inactive Ingredients			
citric acid (UNII: 2968 PHW8 QP) D&C red no. 33 (UNII: 9DBA0 SBB0L) FD&C red no. 40 (UNII: WZB9127XOA) glycerin (UNII: PDC6A3C0OX) methylparaben (UNII: A2I8C7HI9T) propylene glycol (UNII: 6DC9Q167V3) propylparaben (UNII: Z8IX2SC1OH) water (UNII: 059QF0KO0R) sodium benzoate (UNII: OJ245FE5EU)	Ingredient Name	Strength		
D&C red no. 33 (UNII: 9DBA0SBB0L) FD&C red no. 40 (UNII: WZB9127XOA) glycerin (UNII: PDC6A3C0OX) methylparaben (UNII: A2I8C7HI9T) propylene glycol (UNII: 6DC9Q167V3) propylparaben (UNII: Z8IX2SC1OH) water (UNII: 059QF0KO0R) sodium benzoate (UNII: OJ245FE5EU)	alcohol (UNII: 3K9958V90M)			
FD&C red no. 40 (UNII: WZB9127XOA) glycerin (UNII: PDC6A3C0OX) methylparaben (UNII: A2I8C7HI9T) propylene glycol (UNII: 6DC9Q167V3) propylparaben (UNII: Z8IX2SC1OH) water (UNII: 059QF0KO0R) sodium benzoate (UNII: OJ245FE5EU)	citric acid (UNII: 2968PHW8QP)			
glycerin (UNII: PDC6A3C0OX) methylparaben (UNII: A2I8C7HI9T) propylene glycol (UNII: 6DC9Q167V3) propylparaben (UNII: 28IX2SC1OH) water (UNII: 059QF0KO0R) sodium benzoate (UNII: OJ245FE5EU)	D&C red no. 33 (UNII: 9DBA0SBB0L)			
methylparaben (UNII: A2I8C7HI9T) propylene glycol (UNII: 6DC9Q167V3) propylparaben (UNII: Z8IX2SC10H) water (UNII: 059QF0KO0R) sodium benzoate (UNII: OJ245FE5EU)	FD&C red no. 40 (UNII: WZB9127XOA)			
propylene glycol (UNII: 6DC9Q167V3) propylparaben (UNII: Z8IX2SC1OH) water (UNII: 059QF0KO0R) sodium benzoate (UNII: OJ245FE5EU)	glycerin (UNII: PDC6A3C0OX)			
propylparaben (UNII: Z8IX2SC1OH) water (UNII: 059QF0KO0R) sodium benzoate (UNII: OJ245FE5EU)	methylparaben (UNII: A2I8C7HI9T)			
water (UNII: 059QF0KO0R) sodium benzoate (UNII: OJ245FE5EU)	propylene glycol (UNII: 6DC9Q167V3)			
sodium benzoate (UNII: OJ245FE5EU)	propylparaben (UNII: Z8IX2SC1OH)			
	water (UNII: 059QF0KO0R)			
sodium citrate (UNII: 1Q73Q2JULR)	sodium benzoate (UNII: OJ245FE5EU)			
	sodium citrate (UNII: 1Q73Q2JULR)			
sucrose (UNII: C151H8M554)	sucrose (UNII: C151H8M554)			

Product Characteristics			
Color		Score	
Shape		Size	
Flavor	PEPPERMINT (peppermint Flavor)	Imprint Code	
Contains			

Packaging				
Ш	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:54868-6311-0	473 mL in 1 BOTTLE, PLASTIC		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	09/22/2011	

Labeler - Physicians Total Care, Inc. (194123980)

Establishment			
Name	Address	ID/FEI	Business Operations

Physicians Total Care, Inc.	19412398	relabel, repack	

Revised: 6/2011 Physicians Total Care, Inc.