

DOCUSATE SODIUM LIQUID- docusate sodium liquid
LLORENS PHARMACEUTICALS INTERNATIONAL DIVISION

Llorens-Ducosate Sodium 813

Active Ingredients (per 5 mL)

Docusate Sodium 50 mg

Purpose

Stool Softener

Uses

Relief of occasional constipation

Warnings

Do not use when

- abdominal pain, nausea, or vomiting are present unless directed by a doctor
- for more than one week unless directed by a doctor

Ask a doctor before use if you

- are taking mineral oil
- have noticed a sudden change in bowel habits that last more than two weeks

Stop use and ask doctor if

- you have no bowel movements within 3 days
- you have rectal bleeding
- these could be signs of a serious condition
- a skin rash occurs
- you experience throat irritation

If pregnant or breast-feeding, ask a doctor before use

Keep out of reach of children. In case of accidental overdose, seek medical assistance or contact a Poison Control Center right away.

Directions

- follow dosing directions below or use as directed by a physician
- must be given in a 6 oz to 8 oz glass of milk or fruit juice to prevent throat irritation
- may be taken as a single daily dose or in dividend dose
- take maximum dose daily until first bowel movement, dosage should then be reduced according to individual response
- do not exceed recommended dose
- shake well before using

1 teaspoonful = 5 mL

Age	Dose
Adults and children over 12 years of age	1 to 6 teaspoons (50 mg - 300 mg)

Inactive ingredients: FD&C red #40, flavor, methylparaben, poloxamer, polyethylene glycol, propylene glycol, propylparaben, sodium benzoate, sodium citrate, sucralose

Questions or comments? 1-866-595-5598

NDC: 54859-813-16

Drug Facts

Active Ingredient (per 5 mL teaspoonful)	Purpose
Docusate Sodium 50 mg.....	Stool Softener

Uses • Relief of occasional constipation

Warnings
Do not use • if you are currently taking mineral oil, unless directed by a doctor

Ask a doctor before use if you have • stomach pain • nausea • vomiting • notice a sudden change in bowel habits that lasts over 2 weeks

Stop use and ask a doctor if • you have rectal bleeding or fail to have a bowel movement after use. These could be signs of a serious condition • you need to use a stool softener for more than 1 week • a skin rash occurs • you experience throat irritation

If pregnant or breast-feeding, ask a health professional before use.

Keep out of the reach of children. In case of accidental overdose, get medical help or contact Poison Control Center (1-800-222-1222) right away.



DOCUSATE SODIUM LIQUID

50 mg / 5 mL

Drug Facts (continued)

Directions
 • Follow dosing directions below or use as directed by a physician. • Must be given in a 6 oz to 8 oz glass of milk or fruit juice to prevent throat irritation • May be taken as a single daily dose or in divided doses • Take maximum dose daily until first bowel movement, dosage should then be reduced according to individual response • Do not exceed recommended dose • Shake well before using
 1 teaspoonful = 5 mL

Age	Dose
Adults and children over 12 years of age	1 to 6 teaspoons (50 mg- 300 mg)
Children under 12 years of age	Ask a doctor

Other Information:
 • Each teaspoonful (5 mL) contains: sodium 5 mg
 • Protect from excessive heat
 • Protect from freezing
 • Store at room temperature 20-25°C (68-77°F)
 • **Tamper Evident:** Do not use if foil over bottle opening is torn, broken, or missing.

Inactive Ingredients: artificial and natural cherry flavor, FD&C Red #40, methylparaben, poloxamer, polyethylene glycol, propylene glycol, propylparaben, purified water, sodium benzoate, sodium citrate, and sucralose.

Questions or comments: 1-866-595-5598

Manufactured by:
 Llorens Pharmaceutical International Division, Inc.
 7080 NW 37th Ct, Miami, FL 33147

Stool Softener

Lot #:

Exp Date:



Cherry Flavor
16 oz (473 mL)



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Item #: L-63 REV. 08/24



DOCUSATE SODIUM LIQUID

docusate sodium liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54859-813
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DOCUSATE SODIUM (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG)	DOCUSATE SODIUM	50 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	

METHYLPARABEN (UNII: A2I8C7HI9T)	
POLOXAMER 124 (UNII: 1S66E28KXA)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54859-813-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	12/01/2021	

Labeler - LLORENS PHARMACEUTICALS INTERNATIONAL DIVISION (037342305)

Revised: 12/2025

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