

DOCUSATE SODIUM LIQUID- docusate sodium liquid

RUGBY LABORATORIES

Rugby- Ducosate 1304

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, purified water, sodium benzoate, sodium citrate, sucralose

Questions or comments? 1-800-645-2158

Drug Facts

Active ingredient (per 5mL teaspoonful) Purpose
Docusate Sodium 50 mg.....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 a 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 (1-800-222-1222) 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 divided 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



NDC 0536-1304-85

docusate sodium liquid

docusate sodium 50 mg/5 mL stool softener



16 FL OZ (473 mL) cherry flavor

Drug Facts (continued)

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

- **TAMPER-EVIDENT:** Do not use if foil over bottle opening is torn broken or missing.
- store at room temperature 20°C-25°C (68°F-77°F). Permitted excursions between 15°C -30°C (59°F-86°F)
- protect from excessive heat
- Pharmacist-Preserve and dispense in a tight, light-resistant container with a child resistant cap as defined in the USP
- each teaspoon (5 mL) contains sodium 5 mg

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

Questions or comments? (800) 616-2471

THIS IS A BULK CONTAINER NOT INTENDED FOR DISPENSING.

Rev. 05/25 R-164 Re-order No. 371040

Distributed by: RUGBY® LABORATORIES Indianapolis, IN 46268 (800) 616-2471 www.major-rugby.com



Code# L-39

Lot#

Exp. Date:

DOCUSATE SODIUM LIQUID

docusate sodium liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0536-1304
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
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)	

WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0536-1304-85	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	10/01/2020	

Labeler - RUGBY LABORATORIES (079246066)

Revised: 12/2025

RUGBY LABORATORIES