DOCUSATE SODIUM LIQUID- docusate sodium.50mg/5ml solution
Aldama Pharmaceuticals, Inc

Docusate Sodium Liquid

Active ingredient (in each 5 mL-teaspoonful)

Docusate Sodium 50mg

Purpose

Stool Softener

Uses

- Relief of occasional constipation

Warnings

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

Do not use

- abdominal pain, nauseas, 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 withing 3 days
- you have rectal bleeding
- these could be signs of a seriuos 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 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 give in a 6 oz to 8 oz glass of mil 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 the be reduced according to individual response
- do not exceed recommende dose
- shake well before using

Other information

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

Inactive Ingredient

artificial and natural cherry flavor, citric acid, methylparaben, propylene glycol, propylparaben, purified water, sodium citrate, sucralose

Question or Comments?

305-592-9216 or info@aldamapharm.com

DOCUSATE SODIUM LIQUID

docusate sodium.50mg/5ml solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73564-813
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**
SODIUM BENZOATE (UNII: OJ245FE5EU)	
POLOXAMER 124 (UNII: 1S66E28KXA)	
WATER (UNII: 059QF0KO0R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
METHYLPARABEN (UNII: A2I8C7HI9T)	

PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
POLYETHYLENE GLYCOL 500 (UNII: 761NX2Q08Y)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	CHERRY	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73564-813-16	1 in 1 BOX	01/01/2026	
1		473 mL in 1 BOTTLE; 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	01/01/2026	

Labeler - Aldama Pharmaceuticals, Inc (119484030)

Establishment

Name	Address	ID/FEI	Business Operations
Aldama Pharmaceuticals, Inc		119484030	manufacture(73564-813)

Revised: 1/2026 Aldama Pharmaceuticals, Inc