**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 doctore 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

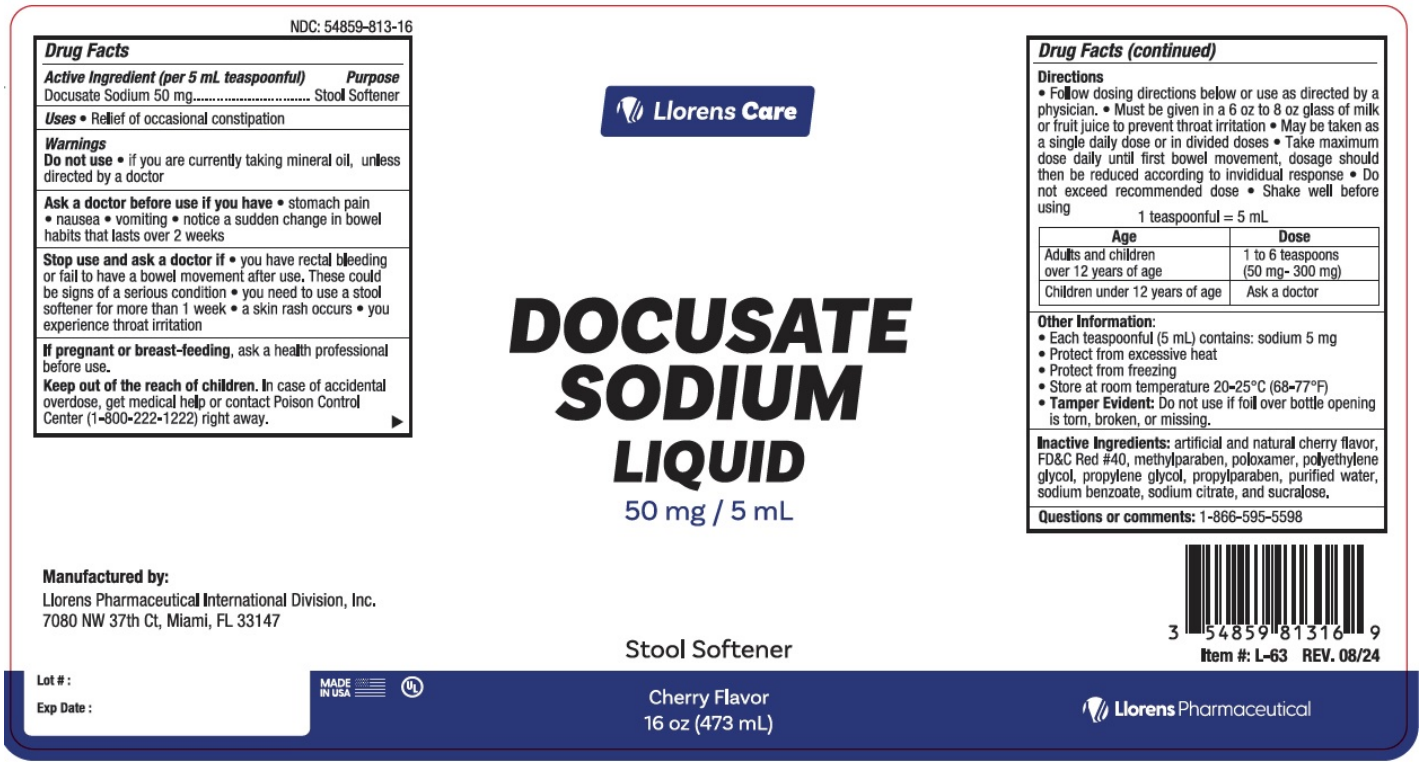
1 teasponnful = 5 mL

| 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 |
|---|---|

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




# 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: 3WQ0SDW1A) | |
| **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 LLORENS PHARMACEUTICALS INTERNATIONAL DIVISION