

**DOCU LIQUID- docusate sodium liquid**  
**Cardinal Health 107, LLC**

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**Drug Facts**

**Active ingredient**

Docusate Sodium 50 mg

**Purpose**

Stool Softener Laxative

**Keep Out of Reach of Children**

**Uses**

- relieves occasional constipation
- generally produces bowel movement in 12-72

**Warnings**

**Do Not Use**

- if you are presently taking mineral oil
- when abdominal pain, nausea, or vomiting are present
- for longer than one week

**Ask a doctor before use if you have**

noticed a sudden change in bowel habits that lasts over two weeks.

**Ask a doctor or pharmacist before use if you are**

taking any other drug. Take this product two or more hours before or after other drugs. Laxatives may affect how other drugs work.

**Stop use and ask a doctor if**

- you have rectal bleeding
- you fail to have a bowel movement after use of this product

These may indicate 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

- may be taken once daily or in divided doses
- give dose in 1/2 glass of milk, fruit juice or infant formula to mask bitter taste and prevent throat irritation

adults and children over 12	1 to 7 teaspoons
children 2 to under 12	1 to 3 teaspoons
children under 2	ask a doctor

## Other information

- **each teaspoon contains:** sodium 5 mg
- shake well before using
- store at controlled room temperature 15° - 30°C (59° - 86°F)
- dispense contents with a child resistant closure in a tight, light resistant container as defined in the USP
- **store in an upright position**

## Inactive Ingredients

D&C Red #33, methylparaben, natural & artificial vanilla flavor, poloxamer 181, polyethylene glycol, propylene glycol, propylparaben, purified water, sodium benzoate. Sodium citrate may be used to adjust pH.

## Questions or comments?

- **Call 1-800-932-5676**

Mon - Thurs. 9:00 am - 4:30 pm EST, Fri. 9:00 am - 2:30 pm EST.

Serious side effects associated with use of this product may be reported to this number.

## Distributed By:

### Cardinal Health

Dublin, OH 43017

L57365410522

## Package/Label Principal Display Panel

DOCU LIQUID

(DOCUSATE SODIUM)

100 mg/ 10 mL

STOOL SOFTENER LAXATIVE

5 Cups



NDC 55154-9630-5

**D108**

**DOCU LIQUID 100 mg / 10 mL**

(Docusate Sodium)

**STOOL SOFTENER LAXATIVE**

**5 CUPS**

Delivers 10 mL

Usual Dosage: See product insert for prescribing information, precautions and warnings.

STORAGE: Store at Controlled Room Temperature  
15 -30 C (59 -86 F)

WARNING: This Unit Dose package is not child resistant  
and is Intended for Institutional Use Only.

KEEP THIS AND ALL DRUGS OUT OF THE REACH  
OF CHILDREN.

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Dublin, OH 43017

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**LOT #: XXXXXXXXXX    EXP. DATE: XX/XX/XX**

## DOCU LIQUID

docusate sodium liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:55154-9630(NDC:50383-349)
<b>Route of Administration</b>	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	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)				
METHYLPARABEN (UNII: A2I8C7HI9T)				
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
PROPYLPARABEN (UNII: Z8IX2SC1OH)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)				
WATER (UNII: 059QF0KO0R)				
POLOXAMER 181 (UNII: 09Y8E6164A)				
Product Characteristics				
Color	PINK		Score	
Shape			Size	
Flavor	VANILLA (natural and artificial flavor)		Imprint Code	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55154-9630-5	5 in 1 BAG	08/01/1997	
1		10 mL in 1 CUP, UNIT-DOSE; 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	08/01/1997	

**Labeler** - Cardinal Health 107, LLC (118546603)