# LACTULOSE- lactulose solution Cardinal Health

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## LACTULOSE SOLUTION, USP

Rx Only

VP2051 04/10

#### **DESCRIPTION**

Lactulose is a synthetic disaccharide in solution form for oral administration. Each 15 mL of lactulose solution contains: 10 g lactulose (and less than 1.6 g galactose, less than 1.2 g lactose, and 1.2 g of other sugars). Also contains FD&C Yellow No. 6, purified water, USP and wild cherry flavoring. A minimal quantity of sodium hydroxide, NF is used to adjust pH when necessary. The pH range is 2.5 to 6.5.

Lactulose is a colonic acidifier which promotes laxation. The chemical name for lactulose is  $4-O-\beta-D$ -galactopyranosyl-D-fructofuranose. It has the following structural formula:

 $C_{12}H_{22}O_{11}$ 

The molecular weight is 342.30. It is freely soluble in water.

#### CLINICAL PHARMACOLOGY

Lactulose is poorly absorbed from the gastrointestinal tract and no enzyme capable of hydrolysis of this disaccharide is present in human gastrointestinal tissue. As a result, oral doses of lactulose reach the colon virtually unchanged. In the colon, lactulose is broken down primarily to lactic acid, and also to small amounts of formic and acetic acids, by the action of colonic bacteria, which results in an increase in osmotic pressure and slight acidification of the colonic contents. This in turn causes an increase in stool water content and softens the stool.

Since lactulose does not exert its effect until it reaches the colon, and since transit time through the colon may be slow, 24 to 48 hours may be required to produce the desired bowel movement.

Lactulose given orally to man and experimental animals resulted in only small amounts reaching the blood. Urinary excretion has been determined to be 3% or less and is essentially complete within 24 hours.

#### INDICATIONS AND USAGE

For the treatment of constipation. In patients with a history of chronic constipation, lactulose solution therapy increases the number of bowel movements per day and the number of days on which bowel movements occur.

#### CONTRAINDICATIONS

Since lactulose solution contains galactose (less than 1.6 g/15 mL), it is contraindicated in patients who require a low galactose diet.

#### WARNINGS

A theoretical hazard may exist for patients being treated with lactulose solution who may be required to undergo electrocautery procedures during proctoscopy or colonoscopy. Accumulation of  $H_2$  gas in significant concentration in the presence of an electrical spark may result in an explosive reaction. Although this complication has not been reported with lactulose, patients on lactulose therapy undergoing such procedures should have a thorough bowel cleansing with a non-fermentable solution. Insufflation of  $CO_2$  as an additional safeguard may be pursued but is considered to be a redundant measure.

#### **PRECAUTIONS**

#### General

Since lactulose solution contains galactose (less than 1.6 g/15 mL) and lactose (less than 1.2 g/15 mL) it should be used with caution in diabetics.

### **Information for Patients**

In the event that an unusual diarrheal condition occurs, contact your physician.

# Laboratory Tests

Elderly, debilitated patients who receive lactulose for more than six months should have serum electrolytes (potassium, chloride, carbon dioxide) measured periodically.

# **Drug Interactions**

Results of preliminary studies in humans and rats suggest that non-absorbable antacids given concurrently with lactulose may inhibit the desired lactulose-induced drop in colonic pH. Therefore, a possible lack of desired effect of treatment should be taken into consideration before such drugs are given concomitantly with lactulose solution.

# Carcinogenesis, Mutagenesis, Impairment of Fertility

There are no known human data on long-term potential for carcinogenicity, mutagenicity, or impairment of fertility.

There are no known animal data on long-term potential for mutagenicity.

Administration of lactulose solution in the diet of mice for 18 months in concentrations of 3 and 10 percent (v/w) did not produce any evidence of carcinogenicity.

In studies of mice, rats, and rabbits, doses of lactulose solution up to 6 or 12 mL/kg/day produced no deleterious effects in breeding, conception, or parturition.

# **Pregnancy**

Teratogenic Effects

Pregnancy category B

Reproduction studies have been performed in mice, rats, and rabbits at doses up to 3 or 6 times the usual human oral dose and have revealed no evidence of impaired fertility or harm to the fetus due to lactulose. There are, however, no adequate and well-controlled studies in pregnant women. Because animal reproduction studies are not always predictive of human response, this drug should be used during pregnancy only if clearly needed.

# **Nursing Mothers**

It is not known whether this drug is excreted in human milk. Because many drugs are excreted in human milk, caution should be exercised when lactulose solution is administered to a nursing woman.

#### **Pediatric Use**

Safety and effectiveness in pediatric patients have not been established.

# **ADVERSE REACTIONS**

Precise frequency data are not available.

Initial dosing may produce flatulence and intestinal cramps, which are usually transient. Excessive dosage can lead to diarrhea with potential complications such as loss of fluids, hypokalemia, and hypernatremia.

Nausea and vomiting have been reported.

#### **OVERDOSAGE**

#### **Signs And Symptoms**

There have been no reports of accidental overdosage. In the event of overdosage, it is expected that diarrhea and abdominal cramps would be the major symptoms. Medication should be terminated.

Oral LD<sub>50</sub>

The acute oral  $LD_{50}$  of the drug is 48.8 mL/kg in mice and greater than 30 mL/kg in rats.

Dialysis

Dialysis data are not available for lactulose. Its molecular similarity to sucrose, however, would suggest that it should be dialyzable.

#### DOSAGE AND ADMINISTRATION

The usual dose is 1 to 2 tablespoonfuls (15 to 30 mL, containing 10 g to 20 g of lactulose) daily. The dose may be increased to 60 mL daily if necessary. Twenty-four to 48 hours may be required to produce a normal bowel movement.

Note: Some patients have found that lactulose solution may be more acceptable when mixed with fruit juice, water or milk.

#### HOW SUPPLIED

Lactulose Solution is available as a cherry-flavored solution, 10 g/15 mL, supplied as follows:

NDC 66689-039-01: 15 mL unit dose cup.

NDC 66689-039-50: Case contains 50 unit dose cups of 15 mL (66689-039-01), packaged in 5 trays of 10 unit dose cups each.

Lactulose Solution is available as a cherry-flavored solution, 20 g/30 mL, supplied as follows:

NDC 66689-038-01: 30 mL unit dose cup.

NDC 66689-038-50: Case contains 50 unit dose cups of 30 mL (66689-038-01), packaged in 5 trays of 10 unit dose cups each.

NDC 66689-038-99: Case contains 100 unit dose cups of 30 mL (66689-038-01), packaged in 10 trays of 10 unit dose cups each.

Lactulose Solution contains lactulose 667 mg/mL (10 g/15 mL).

When ordering this product, include the product code number or NDC in the description.

# Storage

Store at 20°-25°C (68°-77°F) [See USP Controlled Room Temperature]. Avoid subfreezing temperatures.

Under recommended storage conditions, a normal darkening of color may occur. Such darkening is characteristic of sugar solutions and does not affect therapeutic action. Prolonged exposure to temperatures above 86°F (30°C) or to direct light may cause extreme darkening and turbidity which may be pharmaceutically objectionable. If this condition develops, do not use.

Prolonged exposure to freezing temperatures may cause change to a semi-solid, too viscous to pour. Viscosity will return to normal upon warming to room temperature.

Dispense in a tight, light-resistant container as defined in the USP.

**Rx Only.** Keep this and all drugs out of the reach of children.

# Manufactured by:

VistaPharm

Largo, FL 33771

VP2051

04/10

#### PRINCIPAL DISPLAY PANEL

LACTULOSE SOLUTION, USP 20 g/30 mL

5 Cups



T94

# LACTULOSE SOLUTION, USP 20 g/30 mL

5 CUPS

This cup delivers 30 mL

INDICATIONS AND DOSAGE: For the treatment of constipation.

See product insert for prescribing information, precautions and warnings.

STORAGE: Store at 20 - 25 C (68 - 77 F) [See USP CONTROLLED RM TEMP] Avoid subfreezing temperatures.

#### RX ONLY

WARNING: This package is intended for institutional use only. Keep this and all drugs out of the reach of children.

Manufactured by VistaPharm, Inc. Largo, FL 33771

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Repackaged by Cardinal Health Zanesville, OH 43701 L48542120513

LOT#: XXXX

EXP. DATE: 12/34



# **LACTULOSE**

lactulose solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG LABEL	Item Code (Source)	NDC:55154- 4055(NDC:66689-038)
Route of Administration	ORAL	DEA Schedule	

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
LACTULO SE (LACTULO SE)	LACTULOSE	20 g in 30 mL	

Inactive Ingredients		
Ingredient Name	Strength	
FD&C YELLOW NO. 6		
SO DIUM HYDRO XIDE		
WATER		

Product Characteristics			
Color	ORANGE	Score	
Shape		Size	
Flavor	CHERRY	Imprint Code	
Contains			

F	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55154-4055-5	5 in 1 BAG		
1		30 mL in 1 CUP, UNIT-DOSE		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA074138	05/10/2010	

# Labeler - Cardinal Health (188557102)

Establishment				
Name	Address	ID/FEI	Business Operations	
Cardinal Health		188557102	REPACK(55154-4055)	

Revised: 9/2013 Cardinal Health