

LOHIST D- chlorpheniramine maleate / pseudoephedrine hcl liquid
Larken Laboratories, Inc.

LoHist D

Drug Facts

Active Ingredients

(In each 5 mL teaspoonful)

Chlorpheniramine Maleate, USP 2 mg

Pseudoephedrine HCl, USP 30 mg

Purpose

Chlorpheniramine Maleate Antihistamine

Pseudoephedrine HCl Nasal decongestant

Uses

temporarily relieves these symptoms due to hay fever (allergic rhinitis):

- runny nose
- sneezing
- itching of the nose or throat
- itchy, watery eyes
- temporarily relieves nasal congestion due to the common cold, hay fever or other upper respiratory allergies.
- temporarily restores freer breathing through the nose

Warnings

Do not use

- to sedate a child or make a child sleepy
- if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this drug.

Ask a Doctor before use if you have

- heart disease
- high blood pressure
- thyroid disease
- diabetes

- glaucoma
- trouble urinating due to an enlarged prostate gland
- a breathing problem such as emphysema or chronic bronchitis

Ask a doctor or pharmacist before use if you are.

- taking any other nasal decongestant or stimulant
- taking sedatives or tranquilizers

When using this product

Do not exceed recommended dosage.

- marked drowsiness may occur
- avoid alcoholic beverages
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur.
- symptoms do not improve within 7 days or are accompanied by fever.

If pregnant or breast feeding

ask a health professional before use.

Keep out of the reach of children

In case of overdose, get medical help or contact a Poison Control Center immediately.

Directions

Do not exceed 4 doses in a 24-hour period

Age	Dose
Adults and children over 12 years of age	2 teaspoonsful (10 mL) every 4 to 6 hours
Children 6 to under 12 years of age	1 teaspoonful (5 mL) every 4 to 6 hours
Children under 6 years of age	Ask your doctor

Other information

- store at 20°-25°C (68°-77°F)
- very low sodium, contains less than 1 mg sodium per 5 mL teaspoonful

Inactive ingredients

cherry flavoring, methylparaben, polyethylene glycol, propylparaben, purified water saccharin sodium, and sorbitol

Questions or comments?

Call 1-601-855-7678 weekdays from 9:00 am to 4:00 pm CST or go to <http://www.larkenlabs.com>.

Package Label, Principal Display Panel

Figure 1: 16 oz. bottle label

NDC 68047-120-16

LOHIST-D

ANTIHISTAMINE
NASAL DECONGESTANT

SUGAR FREE / ALCOHOL FREE
DYE FREE

Cherry Flavored Liquid

DO NOT USE IF FOIL SEAL UNDER
THE CAP IS BROKEN OR MISSING.

Distributed by:
**LARKEN
LABORATORIES**
Canton, MS 39046

16 fl. oz. (473 mL)

3 68047 12016 8

Lot/Exp. date:

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Rev. 09/2012
400705

PEEL

LOHIST D

chlorpheniramine maleate / pseudoephedrine hcl liquid

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:68047-120

Route of Administration	ORAL
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Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
CHLORPHENIRAMINE MALEATE (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	2 mg in 5 mL
PSEUDOEPHEDRINE HYDROCHLORIDE (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F)	PSEUDOEPHEDRINE HYDROCHLORIDE	30 mg in 5 mL

Inactive Ingredients	
Ingredient Name	Strength
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM ANHYDROUS (UNII: I4807BK602)	
SORBITOL (UNII: 506T60A25R)	

Product Characteristics			
Color		Score	
Shape		Size	
Flavor	CHERRY	Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68047-120-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/15/2003	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	09/15/2003	

Labeler - Larken Laboratories, Inc. (149484540)

Registrant - Larken Laboratories, Inc. (149484540)