NOHIST LQ- chlorpheniramine maleate and phenylephrine hydrochloride liquid
Larken Laboratories, Inc.

NoHist LQ

Drug Facts

Active Ingredients

(In each 5 mL teaspoonful)

Chlorpheniramine Maleate, USP 4 mg

Phenylephrine HCl, USP 10 mg

Purpose

Chlorpheniramine Maleate Antihistamine

Phenylephrine HCl Nasal decongestant

Uses

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

- runny nose
- sneezing
- itchy, watery eyes
- itching of the nose or throat
- 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

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.
- cough or nasal congestion persists for more than 1 week, tends to recur, or is accompanied by a fever, rash or persistent headache. These could be signs of a serious condition.

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 6 doses in a 24-hour period

Age	Dose
Adults and children over 12 years of age	1 teaspoonsful (5 mL) every 4 hours
Children 6 to under 12 years of age	1/2 teaspoonsful (2.5 mL) every 4 hours
Children under 6 years of age	Ask your doctor

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

Inactive Ingredients

bubblegum flavoring, citric acid, D&C Red #33, edetate disodium, glycerin, methylparaben, propylene glycol, propylparaben, purified water, saccharin sodium, and sodium citrate dihydrate

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.

Principal Display Panel

Figure 1: 16 oz. Bottle Label

NOHIST LQ

chlorpheniramine maleate and phenylephrine hydrochloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68047-185
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CHLORPHENIRAMINE MALEATE (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	4 mg in 5 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE -	PHENYLEPHRINE	10 mg

UNII:1WS297W6MV)	HYDROCHLORIDE	in 5 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	

Product Characteristics

Color	pink	**Score**	
Shape		**Size**	
Flavor	BUBBLE GUM	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68047-185-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/06/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	01/06/2011	

Labeler - Larken Laboratories, Inc. (149484540)

Registrant - Larken Laboratories, Inc. (149484540)

Revised: 10/2024 Larken Laboratories, Inc.