

**LOHIST DM- brompheniramine maleate, dextromethorphan hydrobromide and phenylephrine hydrochloride liquid**  
**Larken Laboratories, Inc.**

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**LoHist DM**

***Drug Facts***

**Active ingredients**

***(In each 5 mL teaspoonful)***

Brompheniramine Maleate, USP 2 mg

Dextromethorphan HBr, USP 10 mg

Phenylephrine HCl, USP 5 mg

***Purpose***

Brompheniramine Maleate Antihistamine

Dextromethorphan HBr Antitussive (cough suppressant)

Phenylephrine HCl Nasal decongestant

***Uses***

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

- cough due to minor throat and bronchial irritation
- 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, or associated with sinusitis.
- 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
- a cough that lasts or is chronic such as occurs with smoking, asthma or emphysema
- a cough that occurs with too much phlegm (mucus)

**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	2 teaspoonsful (10 mL) every 4 hours

age

Children 6 to under 12 years of age 1 teaspoonful (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 1 mg sodium per teaspoonful (5 mL)

### ***Inactive Ingredients***

Benzoic acid, edetate disodium, FD&C Red #40, propylene glycol, purified water, saccharin sodium, sorbitol solution, and strawberry flavoring

### ***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**

16 oz Bottle Label

NDC 68047-129-16

# LOHIST·DM

**ANTIHISTAMINE / ANTITUSSIVE  
NASAL DECONGESTANT**

**SUGAR FREE / ALCOHOL FREE**  
**Strawberry 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)**



Lot/Exp. date:

## Drug Facts

### Active ingredients per teaspoonful (5 mL)

	<i>Purpose</i>
Brompheniramine Maleate, USP 2 mg	Antihistamine
Dextromethorphan HBr, USP 10 mg	Antitussive (cough suppressant)
Phenylephrine HCl, USP 5 mg	Nasal decongestant

**Uses** temporarily relieves these symptoms due to hay fever (allergic rhinitis): ■ cough due to minor throat and bronchial irritation ■ 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

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■ 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 two 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

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■ diabetes ■ glaucoma ■ trouble urinating due to an enlarged prostate gland ■ a breathing problem such as emphysema or chronic bronchitis ■ a cough that lasts or is chronic such as occurs with smoking, asthma or emphysema ■ a cough that occurs with too much phlegm (mucus)

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■ taking sedatives or tranquilizers

#### When using this product

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200973-01

**PEEL**

## Drug Facts (continued)

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### Questions or comments?



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## LOHIST DM

brompheniramine maleate, dextromethorphan hydrobromide and phenylephrine hydrochloride liquid

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68047-129
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 5 mL
<b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 5 mL
<b>BROMPHENIRAMINE MALEATE</b> (UNII: IXA7C9ZN03) (BROMPHENIRAMINE - UNII:H57G17P2FN)	BROMPHENIRAMINE MALEATE	2 mg in 5 mL

### Inactive Ingredients

Ingredient Name	Strength
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<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SACCHARIN SODIUM</b> (UNII: SB8ZUX40TY)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>BENZOIC ACID</b> (UNII: 8SKN0B0MIM)	

### Product Characteristics

<b>Color</b>	red	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	STRAWBERRY	<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68047-129-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2012	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	06/01/2012	

**Labeler** - Larken Laboratories, Inc. (149484540)

**Registrant** - Larken Laboratories, Inc. (149484540)