

**BALAMINE DM SYRUP- chlorpheniramine maleate, phenylephrine hydrochloride,
dextromethorphan hydrobromide liquid
Ballay Pharmaceuticals, Inc.**

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Balamine DM Syrup

Drug Facts

Active Ingredients

(in each 5 mL teaspoonful)

Chlorpheniramine Maleate, USP 2 mg

Dextromethorphan HBr, USP 10 mg

Phenylephrine HCl, USP 5 mg

Purposes

Chlorpheniramine Maleate	Antihistamine
Dextromethorphan HBr	Cough Suppressant
Phenylephrine HCl	Nasal Decongestant

▣ *Uses*

temporarily relieves these symptoms due to the common cold, hay fever or other upper respiratory allergies:

- runny nose
- nasal congestion
- sneezing
- itching of the nose or throat
- itchy, watery eyes
- relieves cough associated with the common cold

▣ *Warnings*

▣ *Do not use*

- if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, 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 product.

Ask a doctor before use if you have

- heart disease

- high blood pressure
- thyroid disease
- diabetes
- glaucoma
- a breathing problem, or persistent or chronic cough such as occurs with smoking, asthma, emphysema or chronic bronchitis
- cough accompanied by excessive phlegm (mucus)
- difficulty in urination due to enlargement of the prostate gland.

Ask a doctor or pharmacist before use if you are

□ taking sedatives or tranquilizers.

□ When using this product

- **Do not exceed recommended dosage**
- excitability may occur, especially in children
- marked drowsiness may occur; alcohol, sedatives and tranquilizers may increase the drowsiness effect
- use caution when driving a motor vehicle or operating machinery
- avoid alcoholic beverages.

□ Stop use and ask a doctor if

- new symptoms occur
- nervousness, dizziness, or sleeplessness occur
- cough or nasal congestion persists for more than 7 days, tends to recur, or is accompanied by 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 reach of children.

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

Directions

- **do not exceed 6 doses in a 24 hour period**

adults and children 12 years and over	2 teaspoonsful (10 mL) every 4 hours
children 6 years to under 12 years	1 teaspoonful (5 mL) every 4 hours
children 2 years to under 6 years	1/2 teaspoonful (2.5 mL) every 4 hours
children under 2 years	ask a doctor

□ Other information

- **do not use if tamper evident seal under cap is broken or missing**
- store between 20° - 25° C (68° - 77° F).

□ Inactive Ingredients

citric acid, glycerin, purified water, sodium benzoate, sodium citrate, sorbitol, strawberry flavor.

Questions?

call 1-800-847-1921

Manufactured by:

Ballay Pharmaceuticals, Inc.

Wimberley, Texas 78676

PRINCIPAL DISPLAY PANEL - 473 mL Bottle Label

☐NDC 63162-508-16

Balamine DM

Syrup

Antihistamine/Decongestant/

Antitussive(Cough Suppressant)

Each teaspoonful (5 mL) for oral
administration contains:

Chlorpheniramine

Maleate 2 mg

Dextromethorphan

Hydrobromide10 mg

Phenylephrine

Hydrochloride5 mg

☐For Professional Use Only

Sugar Free/Alcohol Free/

Dye Free

BALLAY 16 fl oz.

(473 mL)

NDC 63162-508-16

Drug Facts

Active ingredients *Purposes*
(in each 5 mL teaspoonful)

Chlorpheniramine Maleate 2 mg Antihistamine
Dextromethorphan HBr 10 mg Cough Suppressant
Phenylephrine HCl 5 mg Nasal Decongestant

Uses temporarily relieves these symptoms due to the common cold, hay fever or other upper respiratory allergies:

- runny nose ■ nasal congestion ■ sneezing
- itching of the nose or throat ■ itchy, watery eyes
- relieves cough associated with the common cold.

Warnings

Do not use

■ if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, 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 product.

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- heart disease ■ high blood pressure
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Balamine DM Syrup

**Antihistamine/Decongestant/
Antitussive(Cough Suppressant)**

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Chlorpheniramine Maleate2 mg
Dextromethorphan Hydrobromide 10 mg
Phenylephrine Hydrochloride5 mg

For Professional Use Only

**Sugar Free/Alcohol Free/
Dye Free**

BALLAY

16 fl. oz.
(473 mL)

Drug Facts (continued)

Stop use and ask a doctor if

- new symptoms occur
- nervousness, dizziness, or sleeplessness occur
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Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

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Manufactured by:
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Wimberley, Texas 78676



Rev. 03/13

BALAMINE DM SYRUP

chlorpheniramine maleate, phenylephrine hydrochloride, dextromethorphan hydrobromide liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63162-508
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLORPHENIRAMINE MALEATE (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	2 mg in 5 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 5 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0K00R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

SODIUM CITRATE (UNI: 1Q73Q2JULR)	
SORBITOL (UNI: 506T60A25R)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	STRAWBERRY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63162-508-16	473 mL in 1 BOTTLE		
2	NDC:63162-508-20	6 in 1 TRAY		
2		20 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	04/23/2013	

Labeler - Ballay Pharmaceuticals, Inc. (035888200)

Establishment

Name	Address	ID/FEI	Business Operations
Ballay Pharmaceuticals, Inc.		035888200	manufacture(63162-508)

Revised: 12/2013

Ballay Pharmaceuticals, Inc.