

**RU-HIST D- brompheniramine maleate and phenylephrine hydrochloride tablet, coated**  
**Allegis Pharmaceuticals, LLC**

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**Ru-Hist D**

**Tablet**

***Drug Facts***

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***Active Ingredients***

<b>Each tablet contains:</b>	<b><i>Purpose</i></b>
Brompheniramine Maleate 4 mg	Antihistamine
Phenylephrine HCl 10 mg	Nasal Decongestant

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**Uses**

Temporarily relieves these symptoms due to the common cold, hay fever (allergic rhinitis) or other upper respiratory allergies:

- runny nose
- sneezing
- itching of the nose or throat
- itchy, watery eyes
- nasal congestion
- reduces swelling of nasal passages

**Warnings**

Do not exceed recommended dosage.

**Do not use this product**

- 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 product

**Ask a doctor before use** if you have

- a breathing problem such as emphysema or chronic bronchitis
- glaucoma
- heart disease
- high blood pressure
- thyroid disease
- diabetes mellitus
- difficulty in urination due to enlargement of the prostate gland

**Do not take this product** if you are taking sedatives or tranquilizers, without first consulting your doctor.

### **When using this product**

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

### **Stop use and ask a doctor if**

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

**If pregnant or breast feeding,** ask a health professional before use.

### **Keep out of reach of children.**

In case of accidental overdose, seek professional help or contact a Poison Control Center immediately.

### **Directions**

#### **Do not exceed recommended dosage.**

Adults and children 12 years of age and over:	1 tablet every 4 hours, not to exceed 6 tablets in 24 hours, or as directed by a doctor
Children 6 to under 12 years of age:	1/2 tablet every 4 hours, not to exceed 3 tablets in 24 hours, or as directed by a doctor
Children under 6 years of age:	Consult a doctor

### **Other information**

Store at 15°-30°C (59°-86°F) [see USP Controlled Room Temperature].

### **Inactive ingredients**

Magnesium Stearate, Microcrystalline Cellulose, Sodium Starch Glycolate

### **Questions or Comments?**

Call 866-633-9033

### **PRINCIPAL DISPLAY PANEL - 60 Tablet Bottle Label**

**NDC 28595-900-60**

**Ru-Hist D**

**Antihistamine • Nasal Decongestant**

**Each tablet contains:**

**Brompheniramine Maleate 4 mg**

**Phenylephrine HCl 10 mg**

**Tamper evident by foil seal under cap. Do not use if foil seal is broken or missing.**

**New Formula**

**ALLEGIS PHARMACEUTICALS**

**Manufactured for:**

**Allegis Pharmaceuticals, LLC**

**Canton, MS 39046**

**60 Tablets**

NDC 28595-900-60

# Ru-Hist D

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**New Formula**



Manufactured for:  
Allegis Pharmaceuticals, LLC  
Canton, MS 39046

60 Tablets

Lot: \_\_\_\_\_  
Exp. Date: \_\_\_\_\_



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Rev. 10/13

NDC 28595-900-60

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**New Formula**

Manufactured for:  
Allegris Pharmaceuticals, LLC  
Canton, MS 39046

**ALLEGRIS PHARMACEUTICALS**

60 Tablets

▲ Lift Here



## RU-HIST D

brompheniramine maleate and phenylephrine hydrochloride tablet, coated

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:28595-900
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>BROMPHENIRAMINE MALEATE</b> (UNII: IXA7C9ZN03) (BROMPHENIRAMINE - UNII:H57G17P2FN)	BROMPHENIRAMINE MALEATE	4 mg
<b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>SODIUM STARCH GLYCOLATE TYPE A CORN</b> (UNII: AG9B65PV6B)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	

**Product Characteristics**

<b>Color</b>	white	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	14mm
<b>Flavor</b>		<b>Imprint Code</b>	300
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:28595-900-60	60 in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2013	

**Marketing Information**

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC Monograph Drug	M012	11/01/2013	

**Labeler** - Allegis Pharmaceuticals, LLC (792272861)