

**ACTIDOM DA- chlorpheniramine maleate, phenylephrine hydrochloride solution**  
**ACTIPHARMA, LLC**

-----  
**ACTIDOM® DA**

***Drug Facts***

**□ Active ingredients (in each 5 mL tsp)**

Chlorpheniramine Maleate 1 mg  
Phenylephrine HCL 2.5 mg

***Purposes***

Antihistamine  
Nasal decongestant

***Uses***

- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies: • runny nose • sneezing • itching of the nose or throat
- itchy, watery eyes • nasal congestion
- temporarily restores free breathing through the nose

***Warnings***

**Do not use** in a child who is 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 child's prescription drug contains an MAOI, ask a doctor or pharmacist before giving this product.

**Ask a doctor before use if child has** • heart disease • diabetes • high blood pressure • thyroid disease • glaucoma • a breathing problem such as chronic bronchitis

**Do not give this product to children who are taking sedatives or tranquilizers, without first consulting the child's doctor.**

**When using this product • do not exceed recommended dosage**

• may cause excitability especially in children • may cause drowsiness • sedatives and tranquilizers may increase the drowsiness effect

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

### ***Directions***

take every 4 hours, or as directed by a doctor.

children 6 to under 12 years of age	take 10 mL (2 tsp). Do not exceed 6 doses in 24 hours
children under 6 years of age	consult a doctor

### ***Other information***

- tamper evident feature: Do not use if safety seal is torn, broken or missing.
- store at controlled room temperature: 15°-30°C (59°-86°F).
- avoid excessive heat or humidity.

### ***Inactive ingredients***

citric acid, D&C Red 33, flavor, hydroxyethyl cellulose, methylparaben, propylene glycol, propylparaben, purified water, sodium citrate, sucralose.

### ***Questions or comments?***

**+1-787-608-0882**

**You may also report serious side effects to this phone number.**

call weekdays from 8AM to 4 PM AST

**Sugar & Alcohol FREE**

**Great Flavor**

Manufactured in USA with imported ingredients for ActiPharma.

San Juan, PR 00917.

[www.actipharma.net](http://www.actipharma.net)

**Packaging**

NDC 63102-115-16

**ACTIDOM<sup>®</sup>  
DA**

**Chlorpheniramine Maleate  
ANTIHISTAMINE  
Phenylephrine HCl  
NASAL DECONGESTANT**

Sugar & Alcohol FREE

Great Flavor



16 fl oz (473 mL)

**Drug Facts**

**Active ingredients (in each 5 mL tsp)**  
Chlorpheniramine Maleate 1 mg.....Antihistamine  
Phenylephrine HCl 2.5 mg.....Nasal decongestant

**Purposes**

**Uses**

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

**Warnings**

**Do not use** in a child who is 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 child's prescription drug contains an MAOI, ask a doctor or pharmacist before giving this product.

**Ask a doctor before use if child has** ■ heart disease ■ diabetes ■ high blood pressure ■ thyroid disease ■ glaucoma ■ a breathing problem such as chronic bronchitis

**Do not give this product to children who are taking sedatives or tranquilizers, without first consulting the child's doctor.**

**When using this product ■ do not exceed recommended dosage**

- may cause excitability especially in children ■ may cause drowsiness ■ sedatives and tranquilizers may increase the drowsiness effect

**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.**

**Drug Facts (continued)**

**Keep out of reach of children.** In case of overdose, get medical help or contact a Poison Control Center right away.

**Directions** ■ take every 4 hours, or as directed by a doctor.

children 6 to under 12 years of age	take 10 mL (2 tsp). Do not exceed 6 doses in 24 hours
children under 6 years of age	consult a doctor

**Other information**

- tamper evident feature: Do not use if safety seal is torn, broken or missing.
- store at controlled room temperature: 15°-30°C (59°-86°F).
- avoid excessive heat or humidity.

**Inactive ingredients**

citric acid, D&C Red 33, flavor, hydroxyethyl cellulose, methylparaben, propylene glycol, propylparaben, purified water, sodium citrate, sucralose.

**Questions or comments?**

+1-877-608-0882

You may also report serious side effects to this phone number. call weekdays from 8AM to 4PM AST

Manufactured in USA with imported ingredients for ActiPharma, San Juan, PR 00917. www.actipharma.net Rev. 04/24



**ACTIDOM DA**

chlorpheniramine maleate, phenylephrine hydrochloride solution

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:63102-115
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>CHLORPHENIRAMINE MALEATE</b> (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	1 mg in 5 mL
<b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	2.5 mg in 5 mL

**Inactive Ingredients**

Ingredient Name	Strength
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	
<b>D&amp;C RED NO. 33</b> (UNII: 9DBA0SBB0L)	
<b>HYDROXYETHYL CELLULOSE, UNSPECIFIED</b> (UNII: T4V6TWG28D)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM CITRATE, UNSPECIFIED FORM</b> (UNII: 1Q73Q2JULR)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	

## Product Characteristics

<b>Color</b>	pink (LIGHT PINK)	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	BANANA, STRAWBERRY	<b>Imprint Code</b>	
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63102-115-16	474 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/20/2022	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	01/20/2022	

**Labeler** - ACTIPHARMA, LLC (079340948)

Revised: 7/2024

ACTIPHARMA, LLC