ALAHIST D- pheniramine maleate, phenylephrine hcl tablet Poly Pharmaceuticals, Inc.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Alahist D

ACTIVE INGREDIENT

Active Ingredient (in each tablet) Purpose

Pheniramine Maleate 17mg	Antihistamine
Phenylephrine HCl 10mg	Nasal Decongestant

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 MA

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 ifyou have

- 🛮 glaucoma
- \sqcap trouble urinating due to enlargement of the prostate gland
- ☐ heart disease
- 🛮 high blood pressure
- 🛮 thyroid disease 🖺
- diabetes

Ask a doctor or pharmacist before use if you are taking sedatives or tranquilizers.

When using this product

- ☐ excitability may occur, especially in children
- 🛮 may cause drowsiness
- □ avoid alcoholic drinks
- 🛘 alcohol, sedatives, and tranquilizers may increase the drowsiness effect
- 🛮 use caution when driving a motor vehicle or operating machinery

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.

DIRECTIONS

Directions

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

INACTIVE INGREDIENTS

Inactive ingredients

Magnesium Stearate, Microcrystalline Cellulose, Natural Yellow, Sodium Starch Glycolate

QUESTIONS

Questions? Comments? Call 1-800-882-1041 Manufactured for: Poly Pharmaceuticals Huntsville, AL 35763 Rev. 04/17

OTHER INFORMATION

Other information

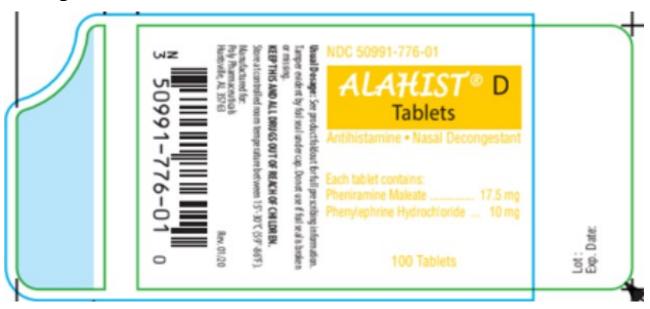
Store at 15°-30°C (59°-86°F). Supplied in a tight, light-resistant container with a child-resistant cap. Alahist D Tablets are yellow, debossed "A" bisect "D" on one side and plain on the other.

Keep out of reach of children

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

Antihistamine

Package Label







ever (allergic rhinitis) or other upper respiratory a nny nose
sneezing itching of the nose or ti



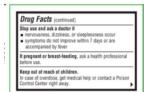




Ask a doctor before use if you have

a breathing problem such as emphysema or chronic bronchilis = glaucoma = trouble urinating due to enlargement of the prostate gland = heart disease = high blood pressure = thyroid disease = diabetes

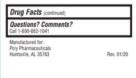








11.8438"



ALAHIST D

pheniramine maleate, phenylephrine hcl tablet

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:50991-776 Route of Administration ORAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
PHENIRAMINE MALEATE (UNII: NYW905655B) (PHENIRAMINE - UNII:134FM9ZZ6M)	PHENIRAMINE MALEATE	17.5 mg	
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg	

Inactive Ingredients			
Ingredient Name	Strength		
MAGNESIUM STEARATE (UNII: 70097M6I30)			
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)			
FERRIC OXIDE YELLOW (UNII: EX43802MRT)			
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6	B)		

Product Characteristics			
Color	yellow	Score	2 pieces
Shape	OVAL	Size	11mm
Flavor		Imprint Code	A;D
Contains			

ı	Packaging				
	# Item Code Package Description		Marketing Start Date	Marketing End Date	
	1	NDC:50991- 776-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	04/01/2020	
	2	NDC:50991- 776-02	12 in 1 BLISTER PACK; Type 0: Not a Combination Product	04/01/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/01/2020	

Labeler - Poly Pharmaceuticals, Inc. (198449894)

Revised: 1/2024 Poly Pharmaceuticals, Inc.