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

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

| 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 MAO |
| 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

- \sqcap a breathing problem such as emphysema or chronic bronchitis
- □ glaucoma
- □ 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

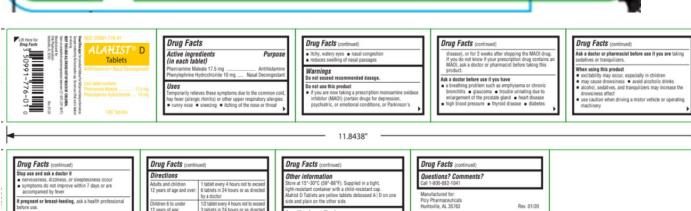
In case of overdose, get medical help or contact a Poison Control Center right away.

Antihistamine

Nasal Decongestant

Package Label





Inactive ingredients

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: 1WS 297W6MV) | PHENYLEPHRINE HYDROCHLORIDE | 10 mg | | |

| In | nactive Ingredients | |
|----|---------------------|----------|
| | Ingredient Name | Strength |

| MAGNESIUM STEARATE (UNII: 70097M6I30) | |
|--|--|
| MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U) | |
| FERRIC OXIDE YELLOW (UNII: EX43802MRT) | |
| SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B) | |

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

| Pa | Packaging Packag | | | | | |
|----|--|---|-------------------------|-----------------------|--|--|
| # | # 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 Application Number or Monograph Category Citation | | Marketing Start Date | Marketing End Date | |
| OTC Monograph Drug | M012 | 04/01/2020 | | |
| | | | | |

Labeler - Poly Pharmaceuticals, Inc. (198449894)

Revised: 7/2024 Poly Pharmaceuticals, Inc.