

STAHIST AD- chlorcyclizine hydrochloride and pseudoephedrine hydrochloride tablet
Magna Pharmaceuticals, Inc.

Stahist AD

Drug Facts

Active Ingredients (in each immediate-release tablet)	Purpose
Chlorcyclizine HCl 25 mg	Antihistamine
Pseudoephedrine HCl 60 mg	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 taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, or emotional conditions, or Parkinson's disease), or for two weeks after stopping the MAOI drug. If you do not know if your prescription drug contains 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 melitus
- difficulty in urination due to enlargement of the prostate gland

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

When using this product

- excitability may occur, especially in children

- may cause drowsiness
- alcohol, sedatives and tranquilizers may increase drowsiness effect
- 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
- 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 by mouth every 6-8 hours, not to exceed 3 tablets in 24 hours, or as directed by a doctor
Children 6 to under 12 years of age:	½ tablet by mouth every 6-8 hours, not to exceed 1½ tablets in 24 hours, or as directed by a doctor
Children under 6 years of age	Consult a doctor

Inactive ingredients

Magnesium Stearate, Microcrystalline Cellulose, Sodium Starch Glycolate

Questions or Comments?

Call 1-888-206-5525 • www.magnaweb.com

Rev. 10/11

58407-0625-30

Stahist AD

Antihistamine • Nasal Decongestant

Each tablet contains:

Chlorcyclizine HCl 25 mg

Pseudoephedrine HCl 60 mg

MAGNA
Pharmaceuticals, Inc.
Louisville, KY 40299

30 Tablets

Add image transcription here...

STAHIST AD

chlorcyclizine hydrochloride and pseudoephedrine hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58407-625
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLORCYCLIZINE HYDROCHLORIDE (UNII: NPB7A7874U) (CHLORCYCLIZINE - UNII:M26C4IP44P)	CHLORCYCLIZINE HYDROCHLORIDE	25 mg
PSEUDOEPHEDRINE HYDROCHLORIDE (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F)	PSEUDOEPHEDRINE HYDROCHLORIDE	60 mg

Inactive Ingredients

Ingredient Name	Strength
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics

Color	white	Score	2 pieces
Shape	OVAL	Size	16mm
Flavor		Imprint Code	M625
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58407-625-30	1 in 1 BOX	12/20/2011	
1		30 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:58407-625-06	6 in 1 BOX	12/20/2011	
2	NDC:58407-625-01	1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:58407-625-01	1 in 1 BLISTER PACK; Type 0: Not a Combination Product	09/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	12/20/2011	

Labeler - Magna Pharmaceuticals, Inc. (620988360)

Registrant - Wittman Pharma, Inc. (830980947)

Revised: 10/2023

Magna Pharmaceuticals, Inc.