AHIST- chlorcyclizine hydrochloride tablet
Magna Pharmaceutcals, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

AHIST™

Drug Facts

Active Ingredients (in each immediate-release tablet)

Chlorcyclizine HCl 25 mg

Purpose

Antihistamine

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

Warnings

Do not exceed recommended dosage.

Do not take this product unless directed by a doctor if you have

- a breathing problem such as emphysema or chronic bronchitis
- glaucoma
- 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
- Use caution when driving a motor vehicle or operating machinery.

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	1 tablet by mouth every 6-8 hours, not to

children 12 years of age and over:	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

Lake Blend Green, Magnesium Stearate, Microcrystalline Cellulose, Sodium Starch Glycolate

Questions or Comments?

Call 1-888-206-5525

www.magnaweb.com

Rev. 12/12

Manufactured for:
MAGNA
Pharmaceuticals, Inc.
Louisville, KY 40299

PRINCIPAL DISPLAY PANEL - 30 Tablet Carton

NDC 58407-025-30

AHIST™

Antihistamine

Each tablet contains:

Chlorcyclizine HCl 25 mg

MAGNA
Pharmaceuticals, Inc.
Louisville, KY 40299

30 Tablets

NDC 58407-025-30

AHIST™

Antihistamine

Each tablet contains:

Chlorcyclizine HCl ..... 25 mg

MAGNA Pharmaceuticals, Inc.
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RETAIN CARTON FOR COMPLETE PRODUCT INFORMATION

Antihistamine

AHIST™

NDC 58407-025-30

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Active Ingredients (in each immediate-release tablet) — **Purpose**
Chlorcyclizine HCl 25 mg ..... Antihistamine

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

Warnings

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■ a breathing problem such as emphysema or chronic bronchitis ■ glaucoma ■ difficulty in urination due to enlargement of the prostate gland

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When using this product
■ excitability may occur, especially in children

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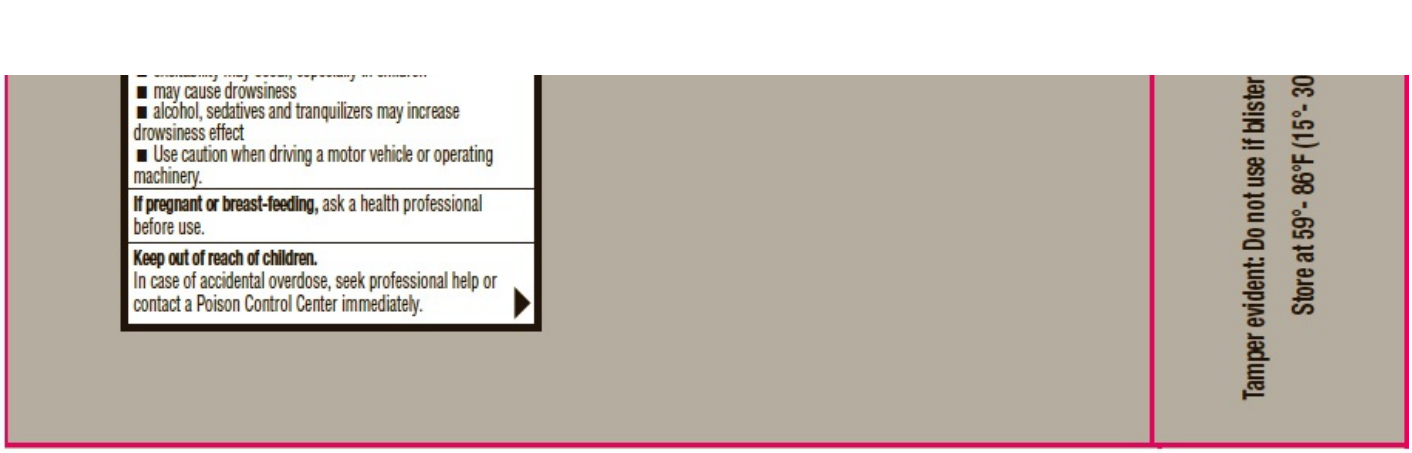
Rev. 12/12

unit is torn, broken or shows any signs of tampering.
°C) [see USP Controlled Room Temperature]



AHIST

chlorcyclizine hydrochloride tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:58407-025
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

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

Product Characteristics			
Color	GREEN	**Score**	2 pieces
Shape	OVAL	**Size**	16mm
Flavor		**Imprint Code**	AHIST;025
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:58407-025-30	30 in 1 BOX		
1	NDC:58407-025-01	1 in 1 BLISTER PACK		
2	NDC:58407-025-06	6 in 1 BOX		
2	NDC:58407-025-01	1 in 1 BLISTER PACK		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part341	01/15/2013	

Labeler - Magna Pharmaceutcals, Inc. (620988360)

Revised: 1/2013 Magna Pharmaceutcals, Inc.