CHLORPHENIRAMINE MALEATE- chlorpheniramine maleate tablet
PD-Rx Pharmaceuticals, Inc.

Active ingredient (in each tablet)

Chlorpheniramine maleate 4 mg

Purpose

Antihistamine

Uses

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

- runny nose
- itchy, watery eyes
- sneezing
- itching of the nose or throat

Warnings

Do not use

to make a child sleepy.

Ask a doctor before use if you have

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

Ask a doctor or pharmacist before use if you are

taking sedatives or tranquilizers.

When using this product

- drowsiness may occur
- avoid alcoholic beverages
- alcohol, sedatives, and tranquilizers may increase drowsiness
- use caution when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

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 (800-222-1222).

Directions

- take every 4 to 6 hours, or as directed by a doctor

adults and children 12 years and over	1 tablet. Do not exceed 6 tablets in 24 hours.
children 6 to under 12 years	1/2 tablet (break tablet in half). Do not exceed 3 whole tablets in 24 hours.
children under 6 years	do not use

Other information

- store at 25ºC (77ºF); excursions permitted between 15º-30ºC (59º-86ºF)
- protect from excessive moisture
- TAMPER EVIDENT: DO NOT USE IF SAFETY SEAL IS BROKEN OR MISSING FROM BOTTLE.

Inactive ingredients

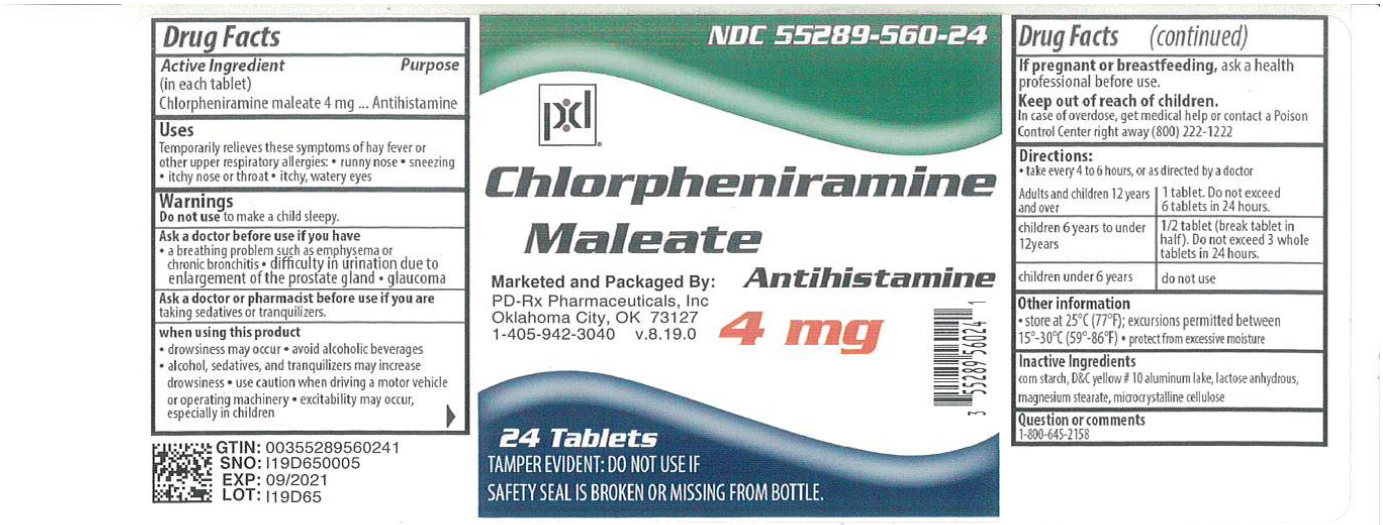
corn starch, D&C yellow #10 aluminum lake, lactose anhydrous, magnesium stearate, microcrystalline cellulose

Questions or comments?

1-800-645-2158

Principal Display Panel

Chlorpheniramine Maleate
4 mg



CHLORPHENIRAMINE MALEATE

chlorpheniramine maleate tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55289-560(NDC:0536-1006)
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CHLORPHENIRAMINE MALEATE (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	4 mg

Inactive Ingredients	
Ingredient Name	**Strength**
STARCH, CORN (UNII: O8232NY3SJ)	
D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6)	
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	

Product Characteristics			
Color	yellow	**Score**	2 pieces
Shape	ROUND	**Size**	8mm

Flavor		**Imprint Code**	44;194
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55289-560-24	24 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/26/2011	
2	NDC:55289-560-30	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/26/2011	
3	NDC:55289-560-01	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/26/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	12/19/1992	

Labeler - PD-Rx Pharmaceuticals, Inc. (156893695)

Registrant - PD-Rx Pharmaceuticals, Inc. (156893695)

Establishment

Name	Address	ID/FEI	Business Operations
PD-Rx Pharmaceuticals, Inc.		156893695	repack(55289-560)

Revised: 1/2025 PD-Rx Pharmaceuticals, Inc.