

CHLORPHENIRAMINE MALEATE 4 MG- chlorpheniramine maleate tablet
PD-Rx Pharmaceuticals, Inc.

Chlorpheniramine Maleate 4 mg

Drug Facts

Active ingredient (in each tablet)

Chlorpheniramine maleate 4 mg

Purpose

Antihistamine

Warnings

Ask a doctor before use if you have

- glaucoma
- trouble urinating due to enlarged prostate gland
- a breathing problem such as emphysema or chronic bronchitis

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

When using this product

- you may get drowsy
- avoid alcoholic drinks
- alcohol, sedatives and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

Drug Facts *continued on back of label*

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.

Directions

adults and children 12 years and over	take 1 tablet every 4 to 6 hours, not more than 6 tablets in 24 hours
children 6 years to under 12 years	take 1/2 tablet every 4 to 6 hours, not more than 3 tablets in 24 hours
children under 6 years	do not use

Other information

- **each tablet contains:** calcium 40 mg
- store at 15°-30°C (59°-86°F)

Inactive ingredients

croscarmellose sodium, D&C yellow# 10 (Al-lake), dicalcium phosphate, magnesium stearate, microcrystalline cellulose, pregelatinized starch, silicon dioxide

Questions or comments?

call **516-341-0666**, 8:30 am - 4:30 pm ET, Monday - Friday

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

* Reliable 1 Laboratories LLC is not affiliated with the owner of the trademark Chlor-Trimeton®

Distributed by: **Reliable 1 Laboratories LLC, Valley Stream, NY 11580**

www.reliable1labs.com

Uses

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

- runny nose
- sneezing
- itchy nose or throat
- itchy, watery eyes

16 HOW SUPPLIED

Chlorpheniramine Maleate Tablets are available as follows:

4 mg: Yellow, round, scored tablet, debossed "heart". Available in bottles of

24 tablets (NDC 72789-134-24)

Chlorpheniramine Maleate 4mg

ANTIHISTAMINE

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Chlorpheniramine maleate 4 mg ...	Antihistamine
Uses	
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If pregnant or breastfeeding, ask a health professional before use.	
	GTIN: 00372789134246 SNO: D21D46000059 EXP: 02/2023 LOT: D21D46

NDC 72789-134-24



Chlorpheniramine Maleate

Antihistamine
4 mg

Marketed and Packaged By:
PD-Rx Pharmaceuticals, Inc
Oklahoma City, OK 73127
1-405-942-3040 v.8.19.0



24 Tablets
TAMPER EVIDENT: DO NOT USE IF
SAFETY SEAL IS BROKEN OR MISSING FROM BOTTLE.

Drug Facts (continued)	
<ul style="list-style-type: none"> • be careful when driving a motor vehicle or operating machinery • excitability may occur, especially in children 	
Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away (800) 222-1222	
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CHLORPHENIRAMINE MALEATE 4 MG

chlorpheniramine maleate tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72789-134(NDC:69618-022)
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

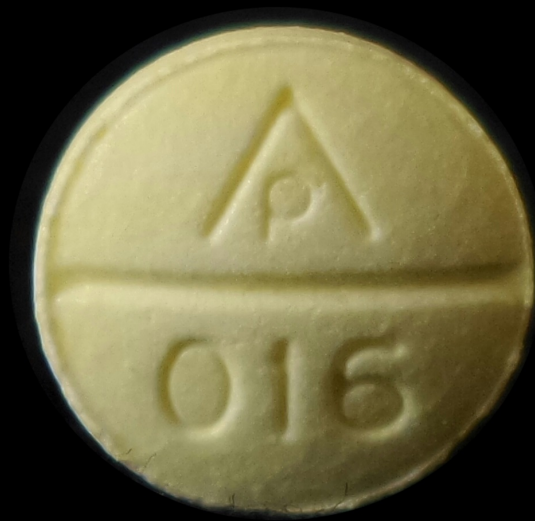
Ingredient Name	Strength
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
CALCIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: L11K75P92J)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
STARCH, PREGELATINIZED CORN (UNII: O8232NY3SJ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

Product Characteristics

Color	yellow	Score	no score
Shape	ROUND	Size	8mm
Flavor		Imprint Code	AP;016
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72789-134-24	24 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/26/2021	



CHLORPHENIRAMINE MALEATE
4mg Tablet

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	11/01/2015	

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(72789-134)

Revised: 10/2024

PD-Rx Pharmaceuticals, Inc.