

**ALLERGY- chlorpheniramine maleate tablet
Bryant Ranch Prepack**

Major 44-194

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
- glaucoma
- difficulty in urination due to enlargement of the prostate gland

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.

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

- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN**
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- protect from excessive moisture
- see end flap for expiration date and lot number

Inactive ingredients

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

Questions or comments?

1-800-426-9391

HOW SUPPLIED

Chlorpheniramine Maleate 4 mg

NDC: 71335-2056-1: 30 Tablets in a BOTTLE, PLASTIC

NDC: 71335-2056-2: 60 Tablets in a BOTTLE, PLASTIC

NDC: 71335-2056-3: 100 Tablets in a BOTTLE, PLASTIC

NDC: 71335-2056-4: 120 Tablets in a BOTTLE, PLASTIC

NDC: 71335-2056-5: 40 Tablets in a BOTTLE, PLASTIC

NDC: 71335-2056-6: 24 Tablets in a BOTTLE, PLASTIC

Repackaged/Relabeled by:

Bryant Ranch Prepack, Inc.
Burbank, CA 91504

Chlorpheniramine Maleate 4 mg Tablet



GTIN 00371335205614
Lot 208620
Exp 4/22/2026
SN 0123456789

Each tablet contains: Chlorpheniramine Maleate, USP 4 mg

Keep this and all drugs out of the reach of children.

Store at 20° to 25° C (68° to 77° F); excursions permitted to 15° to 30° C (59° to 86° F) (see USP controlled Room Temperature).

NDC 71335-2056-1

**Chlorpheniramine Maleate
Tablets**

4 mg

30 Tablets



Repackaged by:
Bryant Ranch Prepack, Inc.
Burbank, CA 91504 USA

Manufactured by:
Major
Pharmaceuticals



ALLERGY

chlorpheniramine maleate tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71335-2056(NDC:0904-0012)
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:71335-2056-1	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/02/2022	
2	NDC:71335-2056-2	60 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/03/2024	
3	NDC:71335-2056-3	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/03/2024	
4	NDC:71335-2056-4	120 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/03/2024	
5	NDC:71335-2056-5	40 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/03/2024	
6	NDC:71335-2056-6	24 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/03/2024	

Marketing Information

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

Labeler - Bryant Ranch Prepack (171714327)

Registrant - Bryant Ranch Prepack (171714327)

Establishment

Name	Address	ID/FEI	Business Operations
Bryant Ranch Prepack		171714327	REPACK(71335-2056) , RELABEL(71335-2056)

Revised: 4/2024

Bryant Ranch Prepack