ALLERGY RELIEF- chlorpheniramine maleate tablet Chain Drug Marketing Association, Inc.

Quality Choice 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
- sneezing
- itchy, watery eyes
- 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.
	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 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

Principal Display Panel

NDC 83324-068-24

QC® QUALITY CHOICE

Allergy Relief

Chlorpheniramine Maleate 4 mg Antihistamine 4 Hour Relief Relieves Sneezing, Runny Nose, Itchy, Watery Eyes, Itchy Throat

Actual Size

24 Tablets

TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING

• SATISFACTION 100% QC® GUARATNTEED •

Distributed by CDMA, Inc. Novi, MI 48375 www.qualitychoice.com Questions: 800-935-2362

50844 REV0419A19408

Questions or comments? 1-800-426-9391

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Drug Facts (continued)

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Buizeeus **■**

- difficulty in unination due to enlargement of the prostate bronchitis 🔳 glaucoma
 - a breathing problem such as emphysema or chronic Ask a doctor before use if you have

Do not use to make a child sleepy. Warnings

- Aching of the nose or throat

 - tever or other upper respiratory allergies:

USBS temporarily relieves these symptoms due to hay

Chlorpheniramine maleate 4 mg. Purpose Active ingredient (in each tablet)

COMPLETE PRODUCT INFORMATION KEEP OUTER PACKAGE FOR

Drug Facts

NDC 83324-068-24



Allergy Relief

Chlorpheniramine Maleate 4 mg

B-0220-194-08-RR REV0419A19408

8 Allergy Relief

Chlorpheniramine Maleate 4 mg



Allergy Relief

Chlorpheniramine Maleate 4 mg

Antihistamine

4 Hour Relief

Relieves Sneezing, Runny Nose, Itchy, Watery Eyes, Itchy Throat

Actual



REV0419A19408

24 Tablets



Distributed by CDMA, Inc. Novi, MI 48375 www.qualitychoice.com Questions: 800-935-2362



Print/No Varnish

Lot & EXP

TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING

Quality Choice 44-194

ALLERGY RELIEF

chlorpheniramine maleate tablet

Pro	duct	Inform	ation
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HUMAN OTC DRUG NDC:83324-068 **Product Type Item Code (Source)**

ORAL **Route of Administration**

Active Ingredient/Active Moiety

Basis of Strength Strength Ingredient Name

CHLORPHENIRAMINE MALEATE (UNII: V1Q0090J9Z) (CHLORPHENIRAMINE -CHLORPHENIRAMINE 4 mg

UNII:3U6IO1965U)

MALEATE

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: 70097M6130)	

MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)

Product Characteristics

Color	yellow	Score	2 pieces
Shape	ROUND	Size	8mm
Flavor		Imprint Code	44;194
Contains			

Packaging

4	tem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324- 068-24	2 in 1 CARTON	09/10/2024	
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
OTC Manager b Deve	MO10	00/10/2024	

OTC Monograph Drug | M012

Labeler - Chain Drug Marketing Association, Inc. (011920774)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(83324-068)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(83324-068) , pack(83324-068)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(83324-068)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(83324-068)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(83324-068)

Revised: 9/2025 Chain Drug Marketing Association, Inc.