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
- 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

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

Principal display panel

QC®
Quality
Choice

NDC 63868-333-24

Allergy Relief
Chlorpheniramine maleate 4 mg
Antihistamine

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

4 Hour Relief

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

50844 ORG041919408

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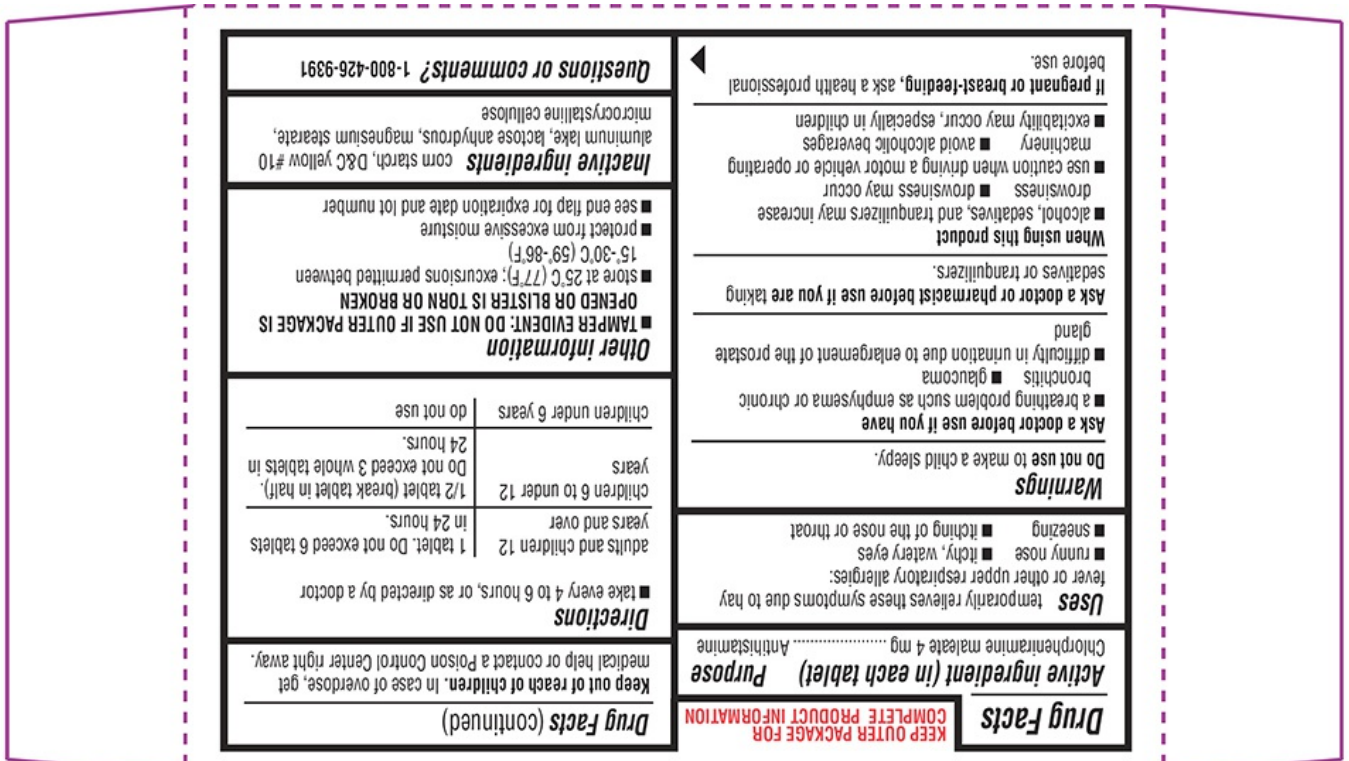
43157 W 9 Mile Rd
Novi, MI 48375
www.qualitychoice.com
Questions: 800-935-2362


Drug Facts
KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION
Active ingredient (in each tablet)
Purpose
Chlorpheniramine maleate 4 mg ..... 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
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When using this product
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Drug Facts (continued)
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Questions or comments? 1-800-426-9391


QC QUALITY CHOICE
Allergy Relief
Chlorpheniramine maleate 4 mg
B-0220-194-08-R
ORG041919408
NDC 63868-333-24
QC QUALITY CHOICE
Allergy Relief
Chlorpheniramine maleate 4 mg
Antihistamine
Relieves Sneezing, Runny Nose, Itchy, Watery Eyes, Itchy Throat
4 Hour Relief
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
No print/No varnish
Lot & Exp date
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Quality Choice 44-194

ALLERGY RELIEF

chlorpheniramine maleate tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-333
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:63868-333-24	2 in 1 CARTON	07/08/2021	
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	07/08/2021	

Labeler - CHAIN DRUG MARKETING ASSOCIATION INC (011920774)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(63868-333)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(63868-333) , pack(63868-333)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(63868-333)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(63868-333)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(63868-333)

Revised: 7/2023 CHAIN DRUG MARKETING ASSOCIATION INC