

CLEMASTINE FUMARATE- clemastine fumarate tablet
Carilion Materials Management

Drug Facts

Active ingredient

Clemastine fumarate 1.34 mg (equivalent to 1 mg clemastine)

Purpose

Antihistamine

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away.

Uses

Temporarily relieves these symptoms of the common cold, hay fever, or other upper respiratory allergies:

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

Warnings

Ask a doctor before use if you have

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

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

If pregnant or breast-feeding,

ask a health professional before use.

Directions

adults and children 12 years and older	1 tablet every 12 hours; not more than 2 tablets in
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adults and children 12 years and older:	24 hours
children under 12 years:	ask a doctor

Other Information

- Safety sealed: do not use if the imprinted bottle seal is open or torn.
- Store at 20°-25°C (68°-77°F).

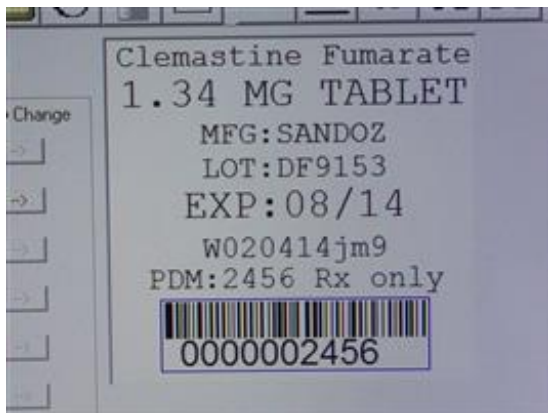
Inactive Ingredients

Hydroxypropyl methylcellulose, lactose (monohydrate), magnesium stearate, microcrystalline cellulose, sodium starch glycolate, and starch (corn).

Questions or comments?

1-800-525-8747

Clemastine Fumarate 1.34 MG TAB



CLEMASTINE FUMARATE			
clemastine fumarate tablet			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68 151-2456(NDC:078 1-1358)
Route of Administration	ORAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
CLEMASTINE FUMARATE (UNII: 19259EGQ3D) (CLEMASTINE - UNII:95QN29S1ID)		CLEMASTINE FUMARATE	1.34 mg
Inactive Ingredients			
Ingredient Name			Strength
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)			
STARCH, CORN (UNII: O8232NY3SJ)			
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D6 1U)			

HYPROMELLOSE 2910 (3 MPAS) (UNII: 0VUT3PMY82)

SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)

MAGNESIUM STEARATE (UNII: 70097M6I30)

Product Characteristics

Color	WHITE	Score	2 pieces
Shape	CAPSULE (capsule shaped)	Size	4mm
Flavor		Imprint Code	GG;159
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68151-2456-0	1 in 1 PACKAGE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA073458	10/31/1993	

Labeler - Carilion Materials Management (079239644)

Registrant - Carilion Materials Management (079239644)

Establishment

Name	Address	ID/FEI	Business Operations
Carilion Materials Management		079239644	REPACK(68151-2456)

Revised: 4/2012

Carilion Materials Management