SUNMARK LORATADINE D- loratadine, pseudoephedrine sulfate tablet, film coated, extended release

Strategic Sourcing Services LLC

McKesson Loratadine D Drug Facts

Active ingredients (in each tablet)

Loratadine 5 mg

Pseudoephedrine sulfate 120 mg

Purpose

Antihistamine

Nasal decongestant

Uses

- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:
- sneezing
- itchy, watery eyes
- runny nose
- itching of the nose or throat
- temporarily relieves nasal congestion due to the common cold, hay fever or other upper respiratory allergies
- reduces swelling of nasal passages
- temporarily relieves sinus congestion and pressure
- temporarily restores freer breathing through the nose

Warnings

Do not use

- if you have ever had an allergic reaction to this product or any of its ingredients
- if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

Ask a doctor before use if you have

- heart disease
- thyroid disease
- high blood pressure
- diabetes
- trouble urinating due to an enlarged prostate gland
- liver or kidney disease. Your doctor should determine if you need a different dose.

When using this product do not take more than directed.

Taking more than directed may cause drowsiness.

Stop use and ask a doctor if

- an allergic reaction to this product occurs. Seek medical help right away.
- symptoms do not improve within 7 days or are accompanied by a fever
- nervousness, dizziness or sleeplessness occurs

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 (1-800-222-1222).

Directions

• do not divide, crush, chew or dissolve the tablet

adults and children 12 years and over	1 tablet every 12 hours; not more than 2 tablets in
J	24 hours
children under 12 years of age	ask a doctor
consumers with liver or kidney disease	ask a doctor

Other information

- each tablet contains: calcium 25 mg
- do not use if blister unit is broken or torn
- store between 20° to 25°C (68° to 77°F)
- keep in a dry place

Inactive ingredients

croscarmellose sodium, dibasic calcium phosphate, hypromellose, lactose monohydrate, magnesium stearate, pharmaceutical ink, povidone, titanium dioxide

Questions or comments?

1-800-719-9260

Package/Label Principal Display Panel

COMPARE TO CLARITIN-D $^{\circledR}$ 12 HOUR EXTENDED RELEASE TABLETS ACTIVE INGREDIENTS

allergy & congestion

12 HOUR

loratadine D

Pseudoephedrine Sulfate 120 mg/Nasal Decongestant

Loratadine 5 mg/Antihistamine

Extended Release Tablets

Relief of:

nasal & sinus congestion due to colds or allergies

sneezing; runny nose; itchy, watery eyes; itchy throat or nose due to allergies

Indoor & Outdoor Allergies

*When taken as directed. See Drug Facts Panel.

NON-DROWSY*

GLUTEN FREE

Actual Size

20 TABLETS





SUNMARK LORATADINE D

loratadine, pseudoephedrine sulfate tablet, film coated, extended release

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70677-0036
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)	LORATADINE	5 mg	
PSEUDO EPHEDRINE SULFATE (UNII: Y9 DL7QPE6B) (PSEUDO EPHEDRINE - UNII:7CUC9 DDI9F)	PSEUDOEPHEDRINE SULFATE	120 mg	

Inactive Ingredients		
Ingredient Name	Strength	
CROSCARMELLOSE SODIUM (UNII: M28 OL1HH48)		
ANHYDRO US DIBASIC CALCIUM PHO SPHATE (UNII: L11K75P92J)		
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)		
LACTO SE MO NO HYDRATE (UNII: EWQ57Q8I5X)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
PO VIDO NE, UNSPECIFIED (UNII: FZ989GH94E)		
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)		

Product Characteristics			
Color	WHITE (to off-white)	Score	no score
Shape	ROUND	Size	12mm
Flavor		Imprint Code	7U0
Contains			

	Packaging			
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:70677-0036-1	20 in 1 CARTON	06/04/2018	
ı	1	1 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
ANDA	ANDA076050	06/04/2018	

Labeler - Strategic Sourcing Services LLC (116956644)

Revised: 10/2020 Strategic Sourcing Services LLC