LORATADINE AND PSEUDOEPHEDRINE- loratadine and pseudoephedrine tablet, film coated, extended release Walgreen Company

Loratadine and Pseudoephedrine

Drug Facts

Active ingredients (in each tablet)	Purpose	
Loratadine, USP 10 mg	Antihistamine	
Pseudoephedrine sulfate, USP 240 mg	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
- reduces swelling of nasal passages
- temporarily relieves sinus congestion and pressure
- temporarily relieves nasal congestion due to the common cold, hay fever or other upper respiratory allergies
- 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

	1 tablet daily with a full glass of water; not more than 1 tablet in 24 hours
children under 12 years of age	ask a doctor
consumers with liver or kidney disease	ask a doctor

Other information

- sodium: contains 10 mg/tablet
- calcium: contains 25 mg/tablet
- TAMPER EVIDENT: DO NOT USE IF BLISTER UNITS ARE TORN, BROKEN OR SHOW ANY SIGNS OF TAMPERING.
- store between 20° C to 25° C (68° F to 77° F)
- protect from light and store in a dry place

Inactive ingredients

calcium carbonate, colloidal silicon dioxide, hydroxypropyl cellulose, hypromellose, iron oxide black, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, povidone, pregelatinized starch, propylene glycol, shellac glaze, sodium alginate, sodium citrate, talc and titanium dioxide

Questions?

call **1-800-406-7984**

DISTRIBUTED BY: WALGREEN CO. 200 WILMOT RD., DEERFIELD, IL 60015

PRINCIPAL DISPLAY PANEL - 240 mg/10 mg Tablet Blister Pack Carton

ORIGINAL PRESCRIPTION STRENGTH NDC 0363-0725-15

Walgreens Compare to the active ingredients in Claritin-D[®] 24 Hour^{††} •WALGREENS• PHARMACIST RECOMMENDED[†]

NON-DROWSY*

Allergy Relief D

Nasal Decongestant

LORATADINE, USP 10 mg / ANTIHISTAMINE PSEUDOEPHEDRINE SULFATE, USP 240 mg / NASAL DECONGESTANT

24 Hour

Indoor & Outdoor Allergies

- Relief of nasal & sinus congestion due to colds or allergies
- Relief of sneezing, runny nose, itchy, watery eyes & itchy throat or nose due to allegies

15 EXTENDED-RELEASE TABLETS

*WHEN TAKEN AS DIRECTED. SEE DRUG FACTS PANEL.

ACTUAL SIZE



Product Inform	nation						
Product Type		HUMAN OTC DRUG	Item Co	ode (Sou	ırce)	NDC:0363	3-0725
Route of Adminis	oute of Administration ORAL						
Active Ingredie	ent/Active	Moiety					
	Ingre	edient Name			Basis of S	Strength	Strengt
LORATADINE (UNII:	7AJO3BO7QN)	(LORATADINE - UNII:7AJO3E	307QN)		LORATADINE		10 mg
PSEUDOEPHEDRIN UNII:7CUC9DDI9F)	E SULFATE (U	NII: Y9DL7QPE6B) (PSEUDO	DEPHEDRIN	NE -	PSEUDOEPHEI SULFATE	DRINE	240 mg
Inactive Ingre	dients						
		Ingredient Name	2			S	trength
CALCIUM CARBON	ATE (UNII: HOG	9379FGK)					
	UNII: ETJ7Z6XB	U4)					
HYDROXYPROPYL	CELLULOSE (1	L600000 WAMW) (UNII: F	RFW2ET671	1P)			
HYPROMELLOSE, U	INSPECIFIED	(UNII: 3NXW29V3WO)					
FERROSOFERRIC C	XIDE (UNII: XM	10M87F357)					
LACTOSE MONOHY	DRATE (UNII:	EWQ57Q8I5X)					
MAGNESIUM STEAI	RATE (UNII: 70	097M6I30)					
MICROCRYSTALLIN	IE CELLULOSI	(UNII: OP1R32D61U)					
POLYETHYLENE GL	YCOL, UNSPE	CIFIED (UNII: 3WJQ0SDW1	A)				
POVIDONE, UNSPE	CIFIED (UNII: I	Z 989GH94E)					
STARCH, CORN (UN	III: 08232NY3S	J)					
PROPYLENE GLYCO	DL (UNII: 6DC9	Q167V3)					
SHELLAC (UNII: 46N	107B71O)						
SODIUM ALGINATE	(UNII: C269C4	G2ZQ)					
SODIUM CITRATE,	UNSPECIFIED	FORM (UNII: 1Q73Q2JULF	R)				
TALC (UNII: 7SEV7J4	R1U)						
TITANIUM DIOXIDE	(UNII: 15FIX9V	2JP)					
Product Chara	cteristics						
Color	white (White	to Off-White)	5	Score		no s	core
Shape	CAPSULE		9	Size		17m	m
Flavor			I	Imprint C	ode	RX72	4
Contains							
Packaging				N#	ting Start	N#	ting End
	Pa			Marko	ting Start	Marka	

	Marketing Category	Application Number or Monograph Citation ANDA076557	Marketing Start Date 11/07/2013	Marketing End Date		
Marketing Information						
3		15 in 1 BLISTER PACK; Type 0: Not a Combination Product				
3	NDC:0363- 0725-15	1 in 1 CARTON	11/07/2013			
2		10 in 1 BLISTER PACK; Type 0: Not a Combination Product				
2	NDC:0363- 0725-69	1 in 1 CARTON	11/07/2013			
1		5 in 1 BLISTER PACK; Type 0: Not a Combination Product				
1	0725-56	1 in 1 CARTON	11/07/2013			

Labeler - Walgreen Company (008965063)

Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		051565745	manufacture(0363-0725)

Revised: 8/2021

Walgreen Company