

PSEUDOEPHEDRINE HYDROCHLORIDE- pseudoephedrine hydrochloride tablet, film coated, extended release

Chain Drug Consortium, LLC

Pseudoephedrine Hydrochloride

Drug Facts

Active ingredient (in each tablet)

Pseudoephedrine HCl, USP 120 mg

Purpose

Nasal decongestant

Uses

- temporarily relieves nasal congestion due to the common cold, hay fever or other upper respiratory allergies
- temporarily relieves sinus congestion and pressure

Warnings

Do not use 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
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland

When using this product do not exceed recommended dosage

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur
- symptoms do not improve within 7 days or occur with a fever

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

adults and children
12 years and over

- take 1 tablet every 12 hours
- do not take more than 2 tablets in 24 hours

children under 12 years	do not use this product in children under 12 years of age
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Other information

- store at 59° to 77° F in a dry place. Protect from light
- **TAMPER EVIDENT: DO NOT USE IF BLISTER UNITS ARE TORN, BROKEN OR SHOW ANY SIGNS OF TAMPERING.**

Inactive ingredients

castor oil, colloidal silicon dioxide, hypromellose, microcrystalline cellulose, magnesium stearate, titanium dioxide

Questions?

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

Distributed By:
Pharmacy Value Alliance, LLC
407 East Lancaster Avenue,
Wayne, PA 19087

PRINCIPAL DISPLAY PANEL - 10 Capsule Blister Pack Carton

†Compare to the active ingredient
in Sudafed® 12 Hour

Premier
Value®

Pseudoephedrine Hydrochloride
Extended-Release Tablets, USP 120 mg

Sinus & Allergy

Long-Acting Nasal Decongestant

Non-Drowsy • 12 Hour

□ SINUS PRESSURE + CONGESTION

□ MAXIMUM STRENGTH

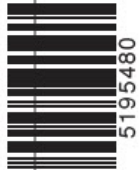
10

Coated Capsule-shaped
Tablets 120 mg Each

INDEPENDENTLY TESTED

PV

SATISFACTION GUARANTEED



5195480



5195480

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Pseudoephedrine Hydrochloride
Extended-Release Tablets, USP 120 mg
Sinus & Allergy
Long-Acting Nasal Decongestant



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✓ SINUS PRESSURE + CONGESTION
✓ MAXIMUM STRENGTH

10 Coated Capsule-shaped
Tablets 120 mg Each



Expiration Date:
Batch No.

NON VARNISH

Keep the carton. It contains important information. See end panel for expiration date.

R0419

†All trademarks are property of their respective owners. This product is not affiliated with the makers/owners of Sudafed® 12 Hour.

Distributed By:
Pharmacy Value Alliance, LLC
407 East Lancaster Avenue,
Wayne, PA 19087
www.emersongroup.com



If for any reason you are not satisfied with this product, please return it to the store where purchased for a full refund.



PSEUDOEPHEDRINE HYDROCHLORIDE

pseudoephedrine hydrochloride tablet, film coated, extended release				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68016-204	
Route of Administration	ORAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
PSEUDOEPHEDRINE HYDROCHLORIDE (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F)		PSEUDOEPHEDRINE HYDROCHLORIDE	120 mg	
Inactive Ingredients				
Ingredient Name			Strength	
CASTOR OIL (UNII: D5340Y2I9G)				
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)				
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				
Product Characteristics				
Color	white	Score	no score	
Shape	CAPSULE	Size	18mm	
Flavor		Imprint Code	204	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016-204-69	10 in 1 BLISTER PACK; Type 0: Not a Combination Product	04/28/2006	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA077442	04/28/2006		

Labeler - Chain Drug Consortium, LLC (101668460)

Registrant - Ohm Laboratories Inc. (184769029)

Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		051565745	MANUFACTURE(68016-204)

Revised: 4/2019

Chain Drug Consortium, LLC