PSEUDOEPHEDRINE HYDROCHLORIDE - pseudoephedrine hydrochloride tablet, extended release
WALGREEN CO.

Pseudoephedrine Hydrochloride Extended-Release Tablets, USP 240 mg

Drug Facts

Active ingredient (in each tablet)

Pseudoephedrine Hydrochloride USP 240 mg

Purpose

Nasal decongestant

Uses

- temporarily relieves nasal congestion due to the common cold, hay fever or other upper respiratory allergies
- reduces swelling of nasal passages
- relieves sinus 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
- had obstruction or narrowing of the bowel Rarely, tablets of this kind may cause bowel obstruction (blockage), usually in people with severe narrowing of the bowel (esophagus, stomach or intestine).

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

- you experience persistent abdominal pain or vomiting

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	• **swallow one** whole tablet with water every 24 hours • **do not exceed one tablet in 24 hours** • **do not divide, crush, chew or dissolve the tablet** • the tablet does not completely dissolve and may be seen in the stool (this is normal)
children under 12 years	do not use this product in children under 12 years of age

Other information

- store at 20° to 25°C (68° to 77°F).
- **do not use if the individual blister unit is open or torn**
- see side panel for lot number and expiration date
- FDA approved dissolution test specifications differ from USP.

Inactive ingredients

black iron oxide, cellulose acetate, colloidal silicon dioxide, dibutyl sebacate, glyceryl mono and dicaprylocaprate, hypromellose, lactose monohydrate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol-part. hydrolyzed, propylene glycol, shellac, sodium lauryl sulfate, stearic acid, talc and titanium dioxide.

Questions or comments?

call **1-855-274-4122** (Monday - Friday 8:30 AM to 5:00 PM EST)

DISTRIBUTED BY: **WALGREEN CO.**
DEERFIELD, IL 60015

MADE IN INDIA

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 1 x 10 Blister Carton Label

Walgreens

NEW

Compare to the active ingredient in Sudafed® Sinus Congestion 24 Hour††

24 HOUR • NON-DROWSY

Nasal Decongestant D

PSEUDOEPHEDRINE HYDROCHLORIDE EXTENDED-RELEASE TABLETS, USP 240 mg / NASAL DECONGESTANT

Maximum Strength 24 Hour
• Sinus pressure & congestion

10
TABLETS ACTUAL SIZE

W00000-0000-0 is a placeholder and must be updated with a unique Packaging Supplier Code issued by GMI for the Walgreens PQA Program. This code is required to be printed on all Walgreens packaging. Additionally, packaging must be produced by a GMI Certified packaging supplier. If the packaging supplier printing the packaging does not have a code they will need to contact GMI to obtain the code and begin the required Certification process by emailing: walgreensmonitoring@sgsco.com

PSEUDOEPHEDRINE HYDROCHLORIDE

pseudoephedrine hydrochloride tablet, extended release

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0363-6611
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PSEUDOEPHEDRINE HYDROCHLORIDE (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F)	PSEUDOEPHEDRINE HYDROCHLORIDE	240 mg

Inactive Ingredients

Ingredient Name	Strength
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
CELLULOSE ACETATE (UNII: 3J2P07GVB6)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
DIBUTYL SEBACATE (UNII: 4W5IH7FLNY)	
GLYCERYL MONO- AND DICAPRYLOCAPRATE (UNII: U72Q2I8C85)	
HYPROMELLOSE 2208 (100000 MPA.S) (UNII: VM7F0B23ZI)	
HYPROMELLOSE 2910 (3 MPA.S) (UNII: 0VUT3PMY82)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SHELLAC (UNII: 46N107B71O)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	WHITE (white to off-white)	**Score**	no score
Shape	ROUND (biconvex)	**Size**	11mm
Flavor		**Imprint Code**	P240
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-6611-10	1 in 1 CARTON	01/30/2026	
1		10 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	ANDA218854	01/30/2026	

Labeler - WALGREEN CO. (008965063)

Registrant - Aurohealth LLC (078728447)

Establishment

Name	Address	ID/FEI	Business Operations
Aurobindo Pharma Limited		650381903	ANALYSIS(0363-6611) , MANUFACTURE(0363-6611)

Revised: 2/2026 WALGREEN CO.