ACTICON- dexbrompheniramine maleate, pseudoephedrine hydrochloride tablet
Actipharma, Inc

ACTICON® COLD & ALLERGY Tablets

Drug Facts

Active ingredients (in each tablet)

Dexbrompheniramine Maleate, USP 2 mg
Pseudoephedrine HCl, USP 60 mg

Purpose

Antihistamine
Nasal Decongestant

Uses

Temporarily relieves these symptoms due to the common cold, hay fever (allergic rhinitis) or other upper respiratory allergies:
• runny nose • sneezing
• itching of the nose or throat
• itchy, watery eyes
• nasal congestion
• reduces swelling of nasal passages

Warnings

Do not exceed recommended dosage.

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
• a breathing problem such as emphysema or chronic bronchitis
• glaucoma • heart disease
• high blood pressure
• thyroid disease
• diabetes mellitus
• difficulty in urination due to enlargement of the prostate gland

Do not take this product if you are taking sedatives or tranquilizers, without

first consulting your doctor.

When using this product
• excitability may occur, especially in children
• may cause drowsiness
• alcohol, sedatives and tranquilizers may increase drowsiness
• avoid alcoholic beverages
• use caution when driving a motor vehicle or operating machinery

Stop use and ask a doctor if
• nervousness, dizziness, or sleeplessness occur
• if symptoms do not improve within 7 days or are accompanied by fever
• new symptoms occur

If pregnant or breast feeding, ask a health professional before use.

Keep out of reach of children. In case of accidental overdose, seek professional help or contact a Poison Control Center immediately.

Directions

Do not exceed recommended dosage.

Adults and children 12 years of age and over:	1 tablet every 4 to 6 hours not to exceed 4 tablets in 24 hours or as directed by a doctor
Children 6 to under 12 years of age:	1/2 tablet every 4 to 6 hours not to exceed 2 tablets in 24 hours or as directed by a doctor
Children under 6 years of age	Consult a doctor

Other information

Store at 15°-30°C (59°-86°F) [see USP Controlled Room Temperature].

Inactive ingredients

Magnesium stearate, microcrystalline cellulose, sodium starch glycolate

Questions or Comments?

call weekdays from 8AM to 4PM AST at **1.787.608.0882**

Contains the same active ingredients as Conex® Tablets*

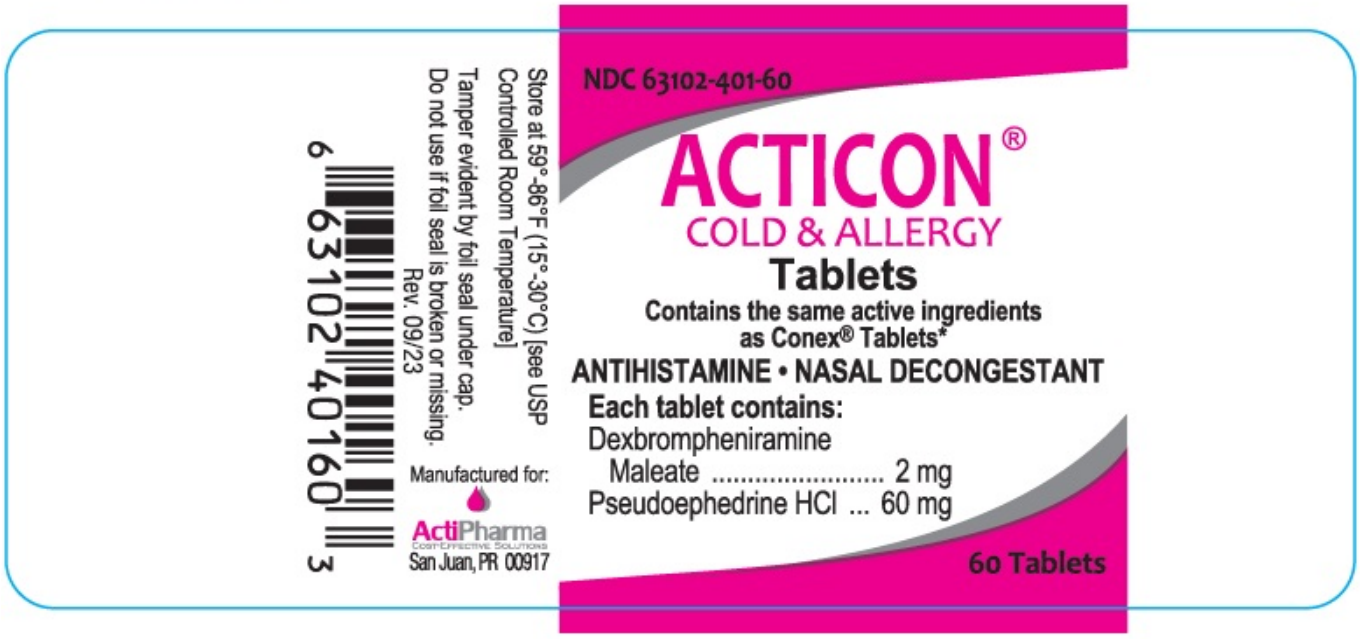
Tamper evident by foil seal under cap.
Do not use if foil seal is broken or missing.

Manufactured in USA with imported ingredients for ActiPharma, Inc. San Juan, PR 00917. www.actipharma.net

***Conex® Tablets is a registered trademark of Llorens Pharmaceutical Corp. This product is not manufactured, distributed or marketed by Llorens Pharmaceutical Corp.**

Packaging

FRONT PANEL



DRUG FACTS TABLE

Drug Facts

Active ingredients (in each tablet)

	Purpose
Dexbrompheniramine Maleate, USP 2 mg	Antihistamine
Pseudoephedrine HCl, USP 60 mg	Nasal Decongestant

Uses Temporarily relieves these symptoms due to the common cold, hay fever (allergic rhinitis) or other upper respiratory allergies:
- runny nose
- sneezing
- itching of the nose or throat

Drug Facts (continued)

In case of accidental overdose, seek professional help or contact a Poison Control Center immediately.

Directions

Do not exceed recommended dosage.

Adults and children 12 years of age and over:	1 tablet every 4 to 6 hours not to exceed 4 tablets in 24 hours or as directed by a doctor
Children 6 to	½ tablet every

- itchy, watery eyes
- nasal congestion
- reduces swelling of nasal passages

Warnings

Do not exceed recommended dosage.

Do not use

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Ask a doctor before use if you have

- a breathing problem such as emphysema or chronic bronchitis
- glaucoma ■ heart disease
- high blood pressure
- thyroid disease
- diabetes mellitus
- difficulty in urination due to enlargement of the prostate gland

Do not take this product if you are taking sedatives or tranquilizers, without first consulting your doctor.

When using this product

- excitability may occur, especially in children
- may cause drowsiness
- alcohol, sedatives and tranquilizers may increase drowsiness
- avoid alcoholic beverages
- use caution when driving a motor vehicle or operating machinery

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur
- if symptoms do not improve within 7 days or are accompanied by fever
- new symptoms occur

If pregnant or breast feeding, ask a health professional before use.
Keep out of reach of children.

Children 6 to under 12 years of age:	½ tablet every 4 to 6 hours not to exceed 2 tablets in 24 hours or as directed by a doctor
Children under 6 years of age	Consult a doctor

Other information

Store at 15°-30°C (59°-86°F) [see USP Controlled Room Temperature].

Inactive ingredients

Magnesium stearate, microcrystalline cellulose, sodium starch glycolate

Questions or Comments?

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Rev. 09/23

NDC 63102-401-60

ACTICON®

COLD & ALLERGY

Tablets

Contains the same active ingredients as Conex® Tablets*

ANTIHISTAMINE • NASAL DECONGESTANT

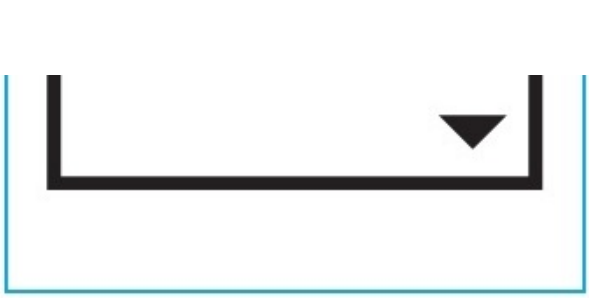
Each tablet contains:
Dexbrompheniramine Maleate ... 2 mg
Pseudoephedrine HCl ..... 60 mg

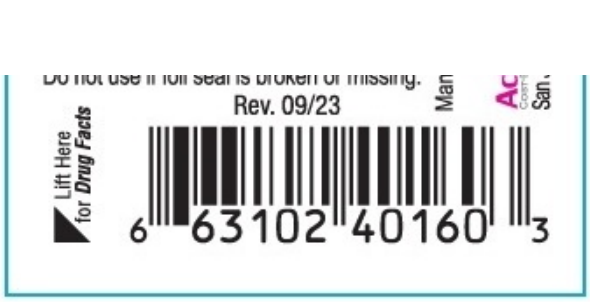
60 Tablets

Store at 59°-86°F (15°-30°C) [see USP Controlled Room Temperature]

Tamper evident by foil seal under cap.
Do not use if foil seal is broken or missing

Manufactured for:
ActiPharma
Effective Solutions
San Juan, PR 00917





ACTICON

dexbrompheniramine maleate, pseudoephedrine hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63102-401
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXBROMPHENIRAMINE MALEATE (UNII: BPA9UT29BS) (DEXBROMPHENIRAMINE - UNII:75T64B71RP)	DEXBROMPHENIRAMINE MALEATE	2 mg
PSEUDOEPHEDRINE HYDROCHLORIDE (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F)	PSEUDOEPHEDRINE HYDROCHLORIDE	60 mg

Inactive Ingredients

Ingredient Name	Strength
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics

Color	white	**Score**	2 pieces
Shape	OVAL	**Size**	13mm
Flavor		**Imprint Code**	A401
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63102-401-60	60 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/13/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	08/13/2015	

Labeler - Actipharma, Inc (079340948)

Revised: 10/2023 Actipharma, Inc