ACTICON- dexbrompheniramine maleate, pseudoephedrine hydrochloride solution
ACTIPHARMA, INC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

NEW ACTICON®*Cold & Allergy Oral Solution*

Drug Facts

Active Ingredients (in each 5 mL tsp)

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:
• relieves sinus congestion and pressure, helps decongest sinus openings and passages
• restores freer breathing through the nose
• runny nose • sneezing • itching of the nose or throat • itchy, watery eyes • nasal congestion

Warnings

Do not exceed recommended dosage
Do not use this product
• 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 • 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 effect • 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 more than 4 doses in 24 hours, or as directed by a doctor.

AGE	***DOSE***
Adults and children 12 years of age and over	1 teaspoonful (5 ml) every 4-6 hours

Other information

• Tamper Evident Feature: Do not use if inner seal is torn, cut, or opened.
• Store at controlled room temperature 15°- 30°C (59°- 86°F).

Inactive ingredients

Citric acid, flavor, methylparaben, potassium citrate, propylene glycol, propylparaben, purified water, sorbitol, sucralose.

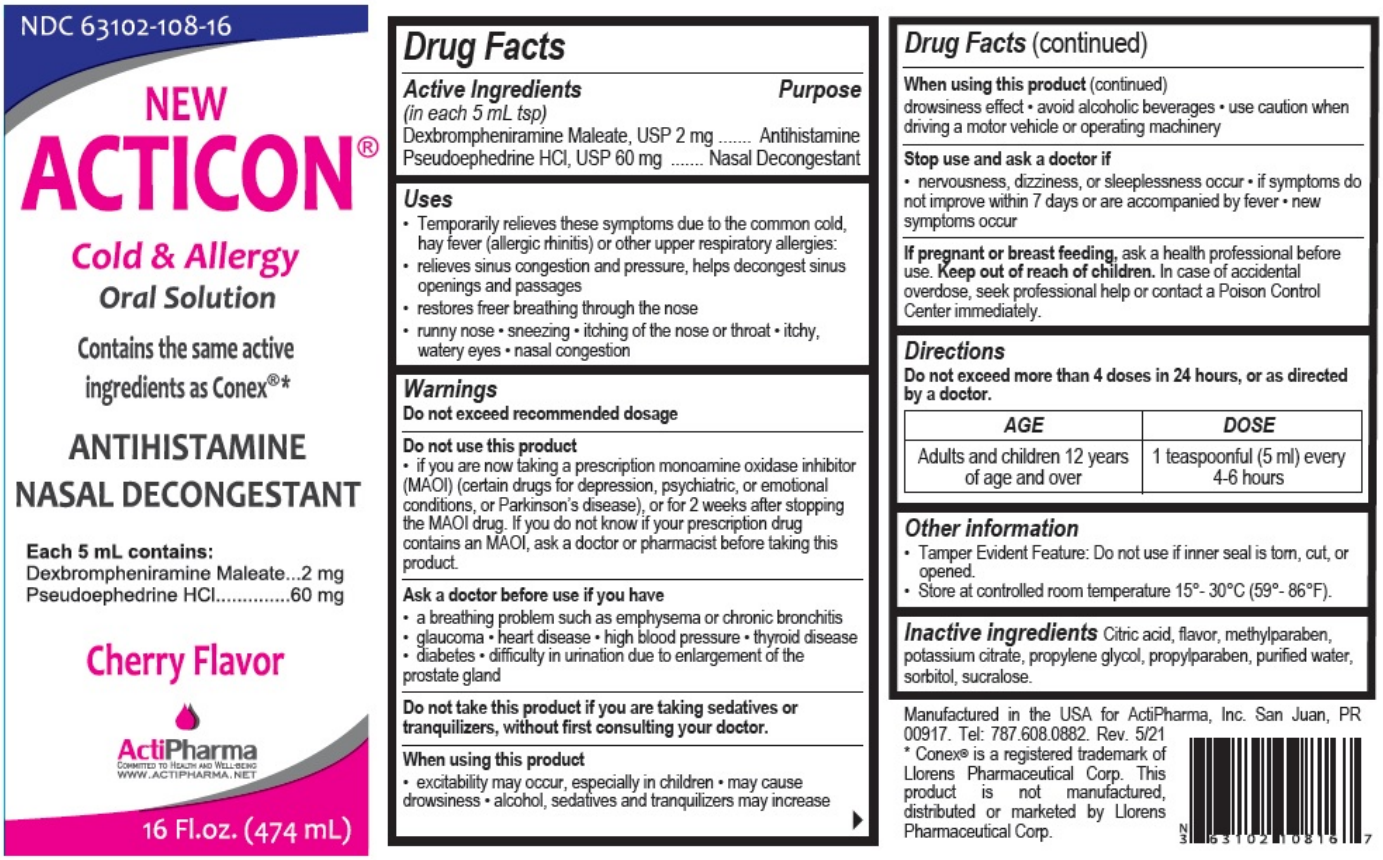
Contains the same active ingredients as Conex®*

Cherry Flavor

ActiPharma
COMMITTED TO HEALTH AND WELL-BEING
WWW.ACTIPHARMA.NET

Manufactured in the USA for ActiPharma, Inc. San Juan, PR 00917. Tel: 787.608.0882.
Rev. 5/21
* Conex® is a registered trademark of Llorens Pharmaceutical Corp. This product is not manufactured, distributed or marketed by Llorens Pharmaceutical Corp.

Packaging



ACTICON

dexbrompheniramine maleate, pseudoephedrine hydrochloride solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63102-108
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 in 5 mL
PSEUDOEPHEDRINE HYDROCHLORIDE (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F)	PSEUDOEPHEDRINE HYDROCHLORIDE	60 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POTASSIUM CITRATE (UNII: EE90ONI6FF)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	

SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics			
Color		**Score**	
Shape		**Size**	
Flavor	CHERRY	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63102-108-16	474 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/15/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	02/15/2022	

Labeler - ACTIPHARMA, INC (079340948)

Revised: 2/2022 ACTIPHARMA, INC