**ACTICON PEDIATRIC- dexbrompheniramine maleate, pseudoephedrine hydrochloride solution**
**ACTIPHARMA, LLC**

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**ACTICON® PEDIATRIC *Cold & Allergy***

***Drug Facts***

***Active ingredients (in each 5 mL tsp)***

Dexbrompheniramine Maleate, USP 1 mg

Pseudoephedrine HCl, USP 30 mg

***Purpose***

Antihistamine

Nasal Decongestant

***Uses***

• Temporarily relieves these symptoms due to the common cold, hay fever (allergic rhinitis) or other 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 • 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 | 2 teaspoonfuls (10 mL) every 4-6 hours |
| Children 6 to under 12 years of age | 1 teaspoonful (5 mL) every 4-6 hours |
| Children under 6 years of age | ask a doctor |

### *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.

### *Questions & Comments?*

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

**Contains the same active ingredients as Conex® Pediatric***

Alcohol FREE

Dye FREE

Sugar FREE

Cherry Flavor

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

Manufactured in the USA with imported ingredients for ActiPharma, LLC. San Juan, PR

00917. *Conex® is a registered trademark of Llorens Pharmaceutical Corp. This product is not manufactured, distributed or marketed by Llorens Pharmaceutical Corp.

## Packaging


# ACTICON PEDIATRIC

dexbrompheniramine maleate, pseudoephedrine hydrochloride solution

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:63102-109 |
| **Route of Administration** | ORAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **DEXBROMPHENIRAMINE MALEATE** (UNII: BPA9UT29BS) (DEXBROMPHENIRAMINE - UNII:75T64B71RP) | DEXBROMPHENIRAMINE MALEATE | 1 mg in 5 mL |
| **PSEUDOEPHEDRINE HYDROCHLORIDE** (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F) | PSEUDOEPHEDRINE HYDROCHLORIDE | 30 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:63102-109-16 | 473 mL in 1 BOTTLE; Type 0: Not a Combination Product | 07/17/2025 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | M012 | 07/17/2025 | |

## Labeler - ACTIPHARMA, LLC (079340948)

Revised: 7/2025 ACTIPHARMA, LLC