WALGREENS CHILDRENS- nighttime cough liquid WALGREENS CO.

Walgreens Children's Night-Time Cough & Chest Congestion

Active ingredients (in each 10 mL)

Chlorpheniramine maleate, USP 2 mg

Dextromethorphan HBr USP 15 mg

Purpose

Antihistamine

Cough suppressant

Uses

- temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold
- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:
- runny nose
- sneezing
- itching of the nose or throat
- itchy, watery eyes

Warnings

Do not us

- to sedate a child or to make a child sleepy
- 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

- trouble urinating due to an enlarges prostate gland
- glaucoma
- a cough that occurs with too much phlegm (mucus)
- a breathing problem or chronic cough that lasts or as occurs with smoking, asthma, chronic bronchitis or emphysema

Ask a doctor or pharmacist before use if you are

taking sedatives or tranquilizers.

When using this product

- do not use more than directed.
- marked drowsiness may occur
- avoid alcoholic drinks
- alcohol, sedatives and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

Stop use and ask a doctor if

• cough last more than 7 days, comes back or is accompanied by fever, rash, or persistent headache. These could be signs of a serious condition.

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 (1-800-222-1222) right away

Directions

- measure only with dosing cup provided.
- keep dosing cup with product
- mL= milliliter
- do not take more than 4 doses in any 24-hour period

age	dose
Children under 6 years	Do not use
Children 6 to under 12 years	10 mL every 6 hours
Adults and children 12 years and older	20 mL every 6 hours

Other information

- each 10 mL contains: sodium 3 mg
- store at room temperature. Do not refrigerate
- contain low sodium
- do not use if printed seal under cap is torn or missing

Inactive ingredients

anhydrous citric acid, FD&C Red # 40, flavor, potassium sorbate, potassium citrate,

propylene glycol, purified water, sodium benzoate, sorbitol, sucralose

Questions or comments?

1-866-467-2748

Principal Display Panel

Compare to the Children's Robitussin $^{\mbox{\scriptsize 8}}$ Nighttime Cough DM active ingredients ††

NDC# 0363-7560-04

Children's

Nighttime Cough

Wal-Tussin® DM

CHLORPHENIRAMINE MALEATE, USP 2 mg/10 mL

ANTIHISTAMINE

DEXTROMETHORPHAN HBr, USP 15 mg/15 mL

COUGH SUPPRESSANT

NIGHTTIME

Relieves cough & runny nose

6 YEARS & OLDER

Fruit Punch Flavor

NATURALLY AND ARTIFICIALLY FLAVORED

Dosage cup included

2-4 FL OZ (118 mL) BOTTLES TOTAL - 8 FL OZ (236 mL)

Walgreens PHARMACIST RECOMMENDED[†]

Health expertise you rely on, quality you trust

[†]Walgreens Pharmacists Survey

*This product is not manufactured or distributed by Pfizer, owner of the registered trademarks Children's Robitussin®

DISTRIBUTED BY: WALGREEN CO.

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Walgreens

100% SATISFACTION GUARANTEED

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TAMPER EVIDENT: DO NOT USE IF PRINTED INNER SEAL UNDER CAP IS BROKEN OR MISSING.

Active ingredients (in each 10 ml):

Chlorpheniramine maleate, USP 2 mg; Dextromethorphan HBr, USP 15 mg

Uses temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold. Temporarily relieves these symptoms due to hay fever or other upper respiratory allergies: runny nose, sneezing, itchy watery eyes, itching of the nose or throat.

Warnings

Do not use to sedate a child or to make a child sleepy or 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.

If pregnant or breast-feeding, ask a health professional before

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away at



4 FL 0Z (118 mL) Artificially Flavored

FRUIT PUNCH FLAVOR

Directions: Do not take more than 4 doses in any 24-hour period. Measure only with dosing cup provided. Keep dosing cup with product, ml = milliliter.

age	dose	
children under 6 years	do not use	
children 6 to under 12 years	10 ml every 6 hours	
adults and children 12 years and older	20 ml every 6 hours	

Other information: each 10 ml contains: sodium 6 mg Store at room temperature. Do not refrigerate. Contains low sodium. Inactive ingredients: anhydrous citric acid, FD&C Red No. 40, flavors, glycerin, lactic acid, potassium sorbate, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol, sucralose.

See carton for full labeling.



WALGREENS CHILDRENS

nighttime cough liquid

Product Information

Product Type HUMAN OTC DRUG NDC:0363-7560 **Item Code (Source)**

• Relieves cough & runny nose

• 6 years & older

Route of Administration ORAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength Strength **CHLORPHENIRAMINE** CHLORPHENIRAMINE MALEATE (UNII: V1Q0090J9Z) (CHLORPHENIRAMINE -2 mg UNII:3U6IO1965U) **MALEATE** in 10 mL DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) **DEXTROMETHORPHAN** 15 ma (DEXTROMETHORPHAN - UNII:7355X3ROTS) **HYDROBROMIDE** in 10 mL

Inactive Ingredients	
Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
LACTIC ACID, UNSPECIFIED FORM (UNII: 33X04XA5AT)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics			
Color	RED	Score	
Shape		Size	
Flavor	FRUIT PUNCH	Imprint Code	
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-7560- 04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/16/2019	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M010	07/16/2019	

Labeler - WALGREENS CO. (008965063)

Revised: 10/2023 WALGREENS CO.