WALGREENS DAYTIME CHILDRENS- cough and chest congestion liquid WALGREENS CO.

Walgreens Children's Cough and Chest Congestion

Active ingredients (in each 5 mL)

Dextromethorphan HBr USP 5 mg Guaifenesin. USP 100 mg

Purposes for Day Time

Cough suppressant Expectorant

Uses

- temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold
- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes

Warnings

Do not us

 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

- cough that occurs with too much phlegm (mucus)
- cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis or emphysema

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 6 doses in any 24-hour period

age	dose	
Children under 4 years Do not use		
Children 4 to under 6 years	5 mL every 4 hours	
Children 6 to under 12 years 10 mL every 4 hours		
Adults and children 12 years and older	20 mL every 4 hours	

Other information

- each 5 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, edetate disodium, FD&C Blue # 1, FD&C Red # 40, flavor, potassium citrate, propylene glycol, purified water, sodium benzoate, sorbitol, sucralose, xanthan gum

Questions or comments?

1-866-467-2748

Principal Display Panel

NDC# 0363-7550-04

Compare to the Children's Robitussin® Daytime Cough & Chest Congestion DM

Children's COUGH & CHEST CONGESTIONS

Wal-Tussin[®] DM

DEXTROMETHORPHAN HBr, USP 5 mg / 5 mL

COUGH SUPPRESSANT

GUAIFENESIN, USP 100 mg / 5 mL

EXPECTORANT

DAYTIME

NON-DROWSY

Relieves cough, chest congestion & mucus

4 YEARS & OLDER

GRAPE FLAVOR

NATURALLY AND ARTIFICIALLY FLAVORED

Dosage cup included

2 - 4 FL OZ (118 mL) BOTTLES

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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WALGREENS DAYTIME CHILDRENS cough and chest congestion liquid **Product Information Product Type** HUMAN OTC DRUG Item Code (Source) ORAL **Route of Administration Active Ingredient/Active Moiety Ingredient Name Basis of Strength** DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) DEXTROMETHORPHAN (DEXTROMETHORPHAN - UNII:7355X3ROTS) HYDROBROMIDE GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ) **GUAIFENES IN Inactive Ingredients Ingredient Name** ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)

Marketing Information

Product

EDETATE DISODIUM (UNII: 7FLD91C86K) FD&C BLUE NO. 1 (UNII: H3R47K3TBD) FD&C RED NO. 40 (UNII: WZB9127XOA) POTASSIUM CITRATE (UNII: EE90ONI6FF) PROPYLENE GLYCOL (UNII: 6DC90167V3)

SODIUM BENZOATE (UNII: OJ245FE5EU)

WATER (UNII: 059QF0K00R)

SORBITOL (UNII: 506T60A25R) SUCRALOSE (UNII: 96K6UQ3ZD4) XANTHAN GUM (UNII: TTV12P4NEE)

Product Characteristics

Color

Shape

Flavor

#

04

Contains

Packaging

Item Code

Marketing

Application Number or Monograph

Package Description

PURPLE

GRAPE

1 NDC:0363-7550- 118 mL in 1 BOTTLE; Type 0: Not a Combination

Score

Size

Imprint Code

Marketing Start

07/16/2019

Marketing Start

Date

Marketing End

Marketing End

Date

NDC:0363-7550

Strength

5 mg

Strength

in 5 mL 100 mg

in 5 mL

Category	Citation	Date	Date
OTC Monograph Drug	M012	07/16/2019	

Labeler - WALGREENS CO. (008965063)

Revised: 10/2023

WALGREENS CO.