WALGREENS CHILDRENS COLD PLUS COUGH PLUS RUNNY NOSEacetaminophen, chlorpheniramine maleate, and dextromethorphan hydrobromide suspension WALGREENS CO.

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Walgreen Children's Cold + Cough + Runny Nose Relief

**Drug Facts** 

Active ingredients (in each 5 Purposes mL)				
Acetaminophen 160 mg	Pain reliever/fever reducer			
Chlorpheniramine maleate 1 mg	Antihistamine			
Dextromethorphan HBr 5 mg	Cough suppressant			

#### Uses

- temporarily relieves the following cold/flu symptoms:
  - minor aches and pains
  - headache
  - sore throat
  - sneezing and runny nose
  - cough
- temporarily reduces fever

# Warnings

# Liver warning

This product contains acetaminophen. Severe liver damage may occur if your child takes

- more than 5 doses in 24 hours, which is the maximum daily amount
- with other drugs containing acetaminophen

**Allergy alert:** acetaminophen may cause severe skin reactions.

Symptoms may include:

- skin reddening
- blisters
- rash

If a skin reaction occurs, stop use and seek medical help right away.

**Sore throat warning:** if sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea, or vomiting, consult a

doctor promptly.

#### Do not use

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- to make a child sleepy
- in a child who is 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 child's prescription drug contains an MAOI, ask a doctor or pharmacist before giving this product.
- if your child has ever had an allergic reaction to this product or any of its ingredients

## Ask a doctor before use if your child has

- liver disease
- a breathing problem such as chronic bronchitis
- glaucoma
- persistent or chronic cough such as occurs with asthma
- cough that occurs with too much phlegm (mucus)

## Ask a doctor or pharmacist before use if your child is

- taking the blood thinning drug warfarin
- taking sedatives or tranquilizers

# When using this product

- do not exceed recommended dose (see overdose warning)
- marked drowsiness may occur
- sedatives and tranquilizers may increase drowsiness
- excitability may occur, especially in children

# Stop use and ask a doctor if

- pain or cough gets worse or lasts more than 5 days
- fever gets worse or lasts more than 3 days
- new symptoms occur
- redness or swelling is present
- cough comes back or occurs with rash or headache that lasts

These could be signs of a serious condition.

# Keep out of reach of children.

# **Overdose warning**

In case of overdose, get medical help or contact a Poison Control Center right away. (1-800-222-1222) Quick medical attention is critical for adults as well as for children even if

you do not notice any signs or symptoms.

#### **Directions**

- this product does not contain directions or complete warnings for adult use
- do not give more than directed (see <u>overdose warning</u>)
- shake well before using
- mL = milliliter
- find right dose on chart below. If possible, use weight to dose; otherwise, use age.
- remove the child protective cap and squeeze your child's dose into the dosing cup
- repeat dose every 4 hours, while symptoms last
- do not give more than 5 times in 24 hours

Weight (lb)	Age (yr)	Dose (mL)
under 36	under 4 years	do not use
36-47	4 to 5 years	do not use unless directed by a doctor
48-95	6 to 11 years	10 mL

**Attention:** use only enclosed dosing cup specifically designed for use with this product. Do not use any other dosing device.

#### Other information

- each 5 mL contains: sodium 3 mg
- store at room temperature

# TAMPER-EVIDENT: DO NOT USE IF PRINTED INNER SEAL UNDER CAP IS TORN OR MISSING

## **Inactive ingredients**

anhydrous citric acid, carboxymethylcellulose sodium, D&C Red No. 33, edetate disodium, FD&C Blue No. 1, FD&C Red No. 40, flavors, glycerin, microcrystalline cellulose, propyl gallate, propylene glycol, purified water, sodium benzoate, sorbitol solution, sucrose, xanthan gum

### Questions or comments?

1-866-467-2748

#### PRINCIPAL DISPLAY PANEL

Compare to the active ingredients in Children's Tylenol® Cold + Cough + Runny Nose\*

NDC 0363-7920-04

Children'sCold + Cough + Runny Nose Relief

**Acetaminophen -** Pain Reliever-Fever Reducer

**Chlorpheniramine Maleate -** Antihistamine **Dextromethorphan HBr -** Cough Suppressant

Oral Suspension

Ages 6-11 Years

- Fever
- Sore Throat
- Sneezing
- Runny Nose
- Cough

## **Grape Flavor**

Naturally and Artificially Flavored

4 FL OZ (120 mL)

IMPORTANT: Keep this carton for future reference on full labeling.

# TAMPER-EVIDENT: DO NOT USE IF PRINTED INNER SEAT UNDER CAP IS TORN OR MISSING.

## Distributed by:

\*This product is not manufactured or distributed by Johnson & Johnson Consumer Inc., distributor of Children's Tylenol® Cold + Cough + Runny Nose.



### WALGREENS CHILDRENS COLD PLUS COUGH PLUS RUNNY NOSE

acetaminophen, chlorpheniramine maleate, and dextromethorphan hydrobromide suspension

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (	Source)	NDC:036	53-7920
Route of Administration	ORAL				
Active Ingredient/Active Moiety					
Ingredient Name			Basis of Stre	ength	Strength
Acetaminophen (UNII: 36209ITL9D) (Acetaminophen - UNII:36209ITL9D)			Acetaminophen		160 mg in 5 mL

<b>Chlorpheniramine Maleate</b> (UNII: V1Q0090J9Z) (Chlorpheniramine - UNII:3U6I01965U)	Chlorpheniramine Maleate	1 mg in 5 mL
<b>Dextromethorphan Hydrobromide</b> (UNII: 9D2RTI9KYH) (Dextromethorphan - UNII:7355X3ROTS)	Dextromethorphan Hydrobromide	5 mg in 5 mL

Inactive Ingredients			
Ingredient Name	Strength		
anhydrous citric acid (UNII: XF417D3PSL)			
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K6790BS311)			
D&C red no. 33 (UNII: 9DBA0SBB0L)			
EDETATE DISODIUM (UNII: 7FLD91C86K)			
FD&C blue no. 1 (UNII: H3R47K3TBD)			
FD&C red no. 40 (UNII: WZB9127XOA)			
glycerin (UNII: PDC6A3C0OX)			
microcrystalline cellulose (UNII: OP1R32D61U)			
PROPYL GALLATE (UNII: 8D4SNN7V92)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
water (UNII: 059QF0KO0R)			
sodium benzoate (UNII: OJ245FE5EU)			
sorbitol (UNII: 506T60A25R)			
sucrose (UNII: C151H8M554)			
xanthan gum (UNII: TTV12P4NEE)			

Product Characteristics				
Color	PURPLE	Score		
Shape		Size		
Flavor	GRAPE	Imprint Code		
Contains				

l	Packaging						
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
	1	NDC:0363- 7920-04	1 in 1 CARTON	02/24/2023			
	1		120 mL in 1 BOTTLE; Type 1: Convenience Kit of Co-Package				

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M013	02/24/2023		

Revised: 11/2024 WALGREENS CO.