

**CHILDRENS TYLENOL COLD PLUS COUGH PLUS RUNNY NOSE- acetaminophen, chlorpheniramine maleate, and dextromethorphan hydrobromide suspension
Johnson & Johnson Consumer Inc.**

Children's TYLENOL[®] COLD + COUGH + RUNNY NOSE

Drug Facts

<i>Active ingredients (in each 5 mL)</i>	<i>Purposes</i>
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 overdose warning)**
- **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

- store between 20-25°C (68-77°F)
- **do not use if carton is opened or if carton tape or bottle wrap imprinted "TYLENOL" is broken or missing**

Inactive ingredients

anhydrous citric acid, D&C red no. 33, FD&C blue no. 1, FD&C red no. 40, flavors, glycerin, microcrystalline cellulose and carboxymethylcellulose sodium, purified water, sodium benzoate, sorbitol solution, sucrose, xanthan gum

Questions or comments?

call **1-800-458-1635** (toll-free) or **215-273-8755** (collect)

PRINCIPAL DISPLAY PANEL

NDC 50580-245-01

**Children's
TYLENOL®**

**COLD + COUGH +
RUNNY NOSE**

Acetaminophen, Pain Reliever-Fever Reducer
Dextromethorphan HBr, Cough Suppressant
Chlorpheniramine Maleate, Antihistamine

Oral Suspension

Ages 6-11 Years

- **FEVER**

- **SORE THROAT**
- **SNEEZING**
- **RUNNY NOSE**
- **COUGH**

4 fl oz (120 mL)

Grape
Flavor



SNAP THIS
CODE OR
TEXT TYKID
TO 87715

MSG & DATA RATES
MAY APPLY
TEXT HELP FOR HELP.
TEXT STOP FOR STOP.



How can we help?
1-800-458-1635

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Oral Suspension

Ages 6-11 Years

- ✓ FEVER
- ✓ SORE THROAT
- ✓ SNEEZING
- ✓ RUNNY NOSE
- ✓ COUGH



Children's **TYLENOL**[®]

COLD + COUGH + RUNNY NOSE

Ages 6-11 Years

• Ibuprofen Free • Alcohol Free • Aspirin Free

DIRECTIONS:

STEP 1.
Shake well
before using



STEP 2.
Remove child
protective
cap



STEP 3.
Squeeze your
child's dose
into the
dosing cup



IMPORTANT:

Please see dosing chart or ask a doctor for your child's correct dose

4 fl oz (120 mL)

Grape
Flavor

PARENTS:

Learn about teen medicine abuse

www.StopMedicineAbuse.org



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Drug Facts (continued)

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Distributed by:

JOHNSON & JOHNSON CONSUMER INC.

McNeil Consumer Healthcare Division

Fort Washington, PA 19034 USA

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www.tylenol.com

30037291



Care To Recycle™

CHILDRENS TYLENOL COLD PLUS COUGH PLUS RUNNY NOSE

acetaminophen, chlorpheniramine maleate, and dextromethorphan hydrobromide suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50580-245
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLORPHENIRAMINE MALEATE (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	1 mg in 5 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	5 mg in 5 mL
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	160 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
D&C RED NO. 33 (UNII: 9DBA05BB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K679OBS311)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBITOL SOLUTION (UNII: 8KW3E207O2)	
SUCROSE (UNII: C151H8M554)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	purple	Score	
Shape		Size	
Flavor	GRAPE	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50580-245-01	1 in 1 CARTON	06/26/2017	
1		120 mL in 1 BOTTLE; Type 1: Convenience Kit of Co-Package		

2	NDC:50580-245-02	1 in 1 CARTON	08/28/2024	
2		240 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	M012		06/26/2017	

Labeler - Johnson & Johnson Consumer Inc. (878046358)

Revised: 4/2024

Johnson & Johnson Consumer Inc.