

**CHLO TUSS- chlophedianol hydrochloride, dexbrompheniramine maleate,  
pseudoephedrine hydrochloride liquid  
R.A. McNeil Company**

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**Chlo Tuss**

**Drug Facts**

**Active ingredients**

**(in each 5 mL teaspoonful)**

Chlophedianol Hydrochloride 12.5 mg

Dexbrompheniramine Maleate 1 mg

Pseudoephedrine Hydrochloride 30 mg

**Purpose**

Cough Suppressant

Antihistamine

Nasal Decongestant

**Uses**

temporarily relieves these symptoms due to the common cold, hay fever (allergic rhinitis) or other upper respiratory allergies:

- cough due to minor throat and bronchial irritation
- runny nose
- sneezing
- itching of the nose or throat
- itchy, watery eyes
- nasal congestion
- reduces swelling of nasal passages

**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 cough that lasts or is chronic such as occurs with smoking, asthma or emphysema
- a cough that occurs with too much phlegm (mucus)
- a breathing problem such as emphysema or chronic bronchitis
- glaucoma
- trouble urinating due to an enlarged prostate gland
- heart disease
- high blood pressure
- thyroid disease
- diabetes

**Ask a doctor or pharmacist before use if you are**

taking sedatives or tranquilizers.

**When using this product**

- excitability may occur, especially in children
- may cause marked drowsiness
- avoid alcoholic drinks
- alcohol, sedatives, and tranquilizers may increase the drowsiness effect
- be careful when driving a motor vehicle or operating machinery

**Stop use and ask a doctor if**

- nervousness, dizziness, or sleeplessness occur
- cough or nasal congestion persists for more than 1 week, tends to recur, or is accompanied by a fever, rash or persistent headache. A persistent cough may be a sign of a serious condition.
- new symptoms occur

**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 right away.

**Directions****Do not exceed recommended dosage.**

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Adults and children 12 years of age and over:	2 teaspoonfuls (10 mL) every 6 hours, not to exceed 8 teaspoonfuls in 24 hours
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Children 6 to under 12 years of age:	1 teaspoonful (5 mL) every 6 hours, not to exceed 4 teaspoonfuls in 24 hours
Children under 6 years of age:	Consult a doctor

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### **Other information**

Store at 59° - 86°F (15° - 30°C)

### **Inactive ingredients**

Citric Acid, Glycerin, Propylene Glycol, Purified Water, Sodium Citrate, Sodium Saccharin, Sorbitol, Tutti Frutti Flavor.

### **Questions? Comments?**

Serious side effects associated with use of this product may be reported to this number. Call 1-423-493-9170  
(8 a.m. to 5 p.m.)

### **PRODUCT PACKAGING**

Tamper evident by foil seal under cap. Do not use if foil seal is broken or missing.

Dispense in a tight, light-resistant container with a child-resistant cap.

**THIS BOTTLE IS NOT TO BE DISPENSED TO THE CONSUMER.**

Manufactured for:  
R.A. McNeil Company  
1150 Latta Street  
Chattanooga, TN 37406-3738

US Patent # 9,050,289

401092-02 Rev. 12/23



NDC 12830-762-16

# Chlo Tuss™

- Cough Suppressant
- Antihistamine
- Nasal Decongestant

## Tutti Frutti Flavor

**EACH 5 mL (1 TEASPOONFUL) CONTAINS:**

Chlophedianol Hydrochloride ..... 12.5 mg  
Dexbrompheniramine Maleate ..... 1 mg  
Pseudoephedrine Hydrochloride ..... 30 mg

**Alcohol Free/Gluten Free  
Sugar Free/Dye Free**

**ONE PINT (473 mL)**

Mfg. for R.A. McNeil Company  
Chattanooga, TN 37406-3738

Rev. 12/23

## Drug Facts ⬆ Lift Here ⬆

### Active ingredients Purpose (in each 5 mL teaspoonful)

Chlophedianol Hydrochloride 12.5 mg .. Cough Suppressant  
Dexbrompheniramine Maleate 1 mg ..... Antihistamine  
Pseudoephedrine Hydrochloride 30 mg ..... Nasal Decongestant

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- runny nose
- sneezing
- itching of the nose or throat
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### Drug Facts (continued)

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Adults and children 12 years of age and over: 2 teaspoonfuls (10 mL) every 6 hours, not to exceed 8 teaspoonfuls in 24 hours

Children 6 to under 12 years of age: 1 teaspoonful (5 mL) every 6 hours, not to exceed 4 teaspoonfuls in 24 hours

Children under 6 years of age: Consult a doctor

### Other information

Store at 59° - 86° F (15° - 30° C)  
Sodium content - 6mg

### Inactive ingredients

Citric Acid, Glycerin, Propylene Glycol, Purified Water, Sodium Citrate, Sodium Saccharin, Sorbitol, Tutti Frutti Flavor.

## CHLO TUSS

chlophedianol hydrochloride, dexbrompheniramine maleate, pseudoephedrine hydrochloride liquid

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:12830-762
<b>Route of Administration</b>	ORAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>CHLOPHEDIANOL HYDROCHLORIDE</b> (UNII: 69QQ58998Y) (CHLOPHEDIANOL - UNII:42C50P12AP)	CHLOPHEDIANOL HYDROCHLORIDE	12.5 mg in 5 mL
<b>DEXBROMPHENIRAMINE MALEATE</b> (UNII: BPA9UT29BS) (DEXBROMPHENIRAMINE - UNII:75T64B71RP)	DEXBROMPHENIRAMINE MALEATE	1 mg in 5 mL
<b>PSEUDOEPHEDRINE HYDROCHLORIDE</b> (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F)	PSEUDOEPHEDRINE HYDROCHLORIDE	30 mg in 5 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM CITRATE</b> (UNII: 1Q73Q2JULR)	
<b>SACCHARIN SODIUM</b> (UNII: SB8ZUX40TY)	
<b>SORBITOL</b> (UNII: 506T60A25R)	

## Product Characteristics

<b>Color</b>		<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	TUTTI FRUTTI	<b>Imprint Code</b>	
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:12830-762-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2013	
2	NDC:12830-762-15	15 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/09/2015	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	10/01/2013	

Revised: 10/2025

R.A. McNeil Company