

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

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.**

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

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 ⬆

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Dexbrompheniramine Maleate 1 mg Antihistamine
Pseudoephedrine Hydrochloride 30 mg Nasal Decongestant

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Other information

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Sodium content - 6mg

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CHLO TUSS

chlophedianol hydrochloride, dexbrompheniramine maleate, pseudoephedrine hydrochloride liquid

Product Information

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

Active Ingredient/Active Moiety

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

Inactive Ingredients

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

Product Characteristics

Color		Score	
Shape		Size	
Flavor	TUTTI FRUTTI	Imprint Code	
Contains			

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: 3/2024

R.A. McNeil Company