

VANACOF CP ALLERGY / COUGH- chlophedianol hcl, pyrilamine maleate solution
GM Pharmaceuticals, INC

VANACOF CP Allergy / Cough

Active Ingredients (in each 15 mL (1TBSP))

Chlophedianol HCl 12.5 mg

Pyrilamine Maleate 25 mg

Purpose

Cough Suppressant

Antihistamine

Uses

- temporarily relieves these symptoms due to the common cold, hay fever or other upper respiratory allergies:
- runny nose ■ itching of the nose or throat
- sneezing ■ itchy, watery eyes
- cough due to minor throat and bronchial irritation

Warnings

Ask a doctor before use if you have

- a cough that lasts or is chronic such as occurs with smoking, asthma, or emphysema
- glaucoma
- difficulty in urination due to enlargement of the prostate gland
- a cough that occurs with too much phlegm (mucus)
- a breathing problem such as emphysema or chronic bronchitis

Ask a doctor before use if you are

taking sedatives or tranquilizers

When using this product

- do not use more than directed ■ avoid alcoholic drinks
- marked drowsiness may occur
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occurs
- symptoms do not improve within 7 days, tend to recur, or are 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 take more than 4 doses in any 24-hour period
- dose as follows or as directed by a doctor ■ mL= milliliter

adults and children 12 years of age and over:	30 mL (2 TBSP) every 6 to 8 hours, not to exceed 120 mL (8 TBSP) per 24 hours
children 6 to under 12 years of age:	15 mL (1 TBSP) every 6 to 8 hours, not to exceed 60 mL (4 TBSP) per 24 hours
children under 6 years:	consult a doctor.

Other information

- store at 20° to 30°C (68° to 86°F)
- each 15 mL (1 TBSP) contains: **Sodium 8.5 mg**

Inactive ingredients

citric acid anhydrous, glycerin, propylene glycol, purified water, sodium benzoate, sodium citrate, sodium saccharin, sorbitol

Questions?

1-888-535-0305 9 a.m. - 5 p.m. CST

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

U.S. Pat: 9,463,191

Distributed by: GM Pharmaceuticals, Inc. Arlington, TX 76018

NDC 58809-189-08

VANACOF® CP

Allergy / Cough

Each 15 mL (1 TBSP) contains:

Chlophedianol HCl 12.5 mg

Pyrilamine Maleate 25 mg

Cough Suppressant • Antihistamine

Alcohol Free / Sugar Free / Gluten Free / Dye Free

GM Pharmaceuticals, Inc.

8 fl oz (237 mL)

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Ask a doctor before use if you have <ul style="list-style-type: none"> ■ a cough that lasts or is chronic such as occurs with smoking, asthma, or emphysema ■ glaucoma ■ difficulty in urination due to enlargement of the prostate gland ■ a cough that occurs with too much phlegm (mucus) ■ a breathing problem such as emphysema or chronic bronchitis 	
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<ul style="list-style-type: none"> ■ do not use more than directed ■ marked drowsiness may occur ■ avoid alcoholic drinks ■ alcohol, sedatives, and tranquilizers may increase drowsiness ■ be careful when driving a motor vehicle or operating machinery ■ excitability may occur, especially in children 	

Drug Facts (continued)							
Stop use and ask a doctor if							
<ul style="list-style-type: none"> ■ nervousness, dizziness, or sleeplessness occurs ■ symptoms do not improve within 7 days, tend to recur, or are 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.							
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VANACOF CP ALLERGY / COUGH

chlophedianol hcl, pyrilamine maleate solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58809-189
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 15 mL
PYRILAMINE MALEATE (UNII: R35D29L3ZA) (PYRILAMINE - UNII:HPE317O9TL)	PYRILAMINE MALEATE	25 mg in 15 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58809-189-08	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/06/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	12/06/2023	

Labeler - GM Pharmaceuticals, INC (793000860)

Revised: 10/2025

GM Pharmaceuticals, INC