

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

VANACOF CP Allergy / Cough

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

Chlophedianol HCl 12.5 mg

Pyrillamine 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. Fort Worth, TX 76118

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)



8 fl oz (237 mL)

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

Drug Facts

Active ingredients (in each 15 mL (1TBSP))
 Chlophedianol HCl 12.5 mg.....Cough Suppressant
 Pyrilamine Maleate 25 mg.....Antihistamine

Purpose

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

Drug Facts (continued)

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

Distributed by: GM Pharmaceuticals, Inc. Fort Worth, TX 76118

U.S. Pat: 9,463,191



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: 12/2023

GM Pharmaceuticals, INC