**VANACOF- chlophedianol hydrochloride, dexchlorpheniramine maleate, and pseudoephedrine hydrochloride liquid**
**GM Pharmaceuticals, INC**

---

**VanaCof ®**

***Drug Facts***

***Active ingredients (in each 5 mL teaspoonful)***

Chlophedianol Hydrochloride 12.5 mg

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

***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
- trouble urinating due to an enlarged prostate gland
- glaucoma
- heart disease
- high blood pressure
- thyroid disease
- diabetes

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

taking sedatives or tranquilizers.

**When using this product**
- excitability may occur, especially in children
- may cause marked drowsiness
- alcohol, sedatives, and tranquilizers may increase the drowsiness effect
- avoid alcoholic drinks
- use caution 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 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 every 6 hours, not to exceed 8 teaspoonfuls in 24 hours. |
| children 6 to under 12 years of age: | 1 teaspoonful every 6 hours, not to exceed 4 teaspoonfuls in 24 hours. |
| children under 6 years of age: | consult a doctor. |

***Other information***
- this package is child resistant

- store at room temperature of 68°-86°F (20°-30°C) with excursions of 59°-86°F (15°-30°C)

### *Inactive ingredients*

Citric acid anhydrous, flavor, glycerin, propylene glycol, purified water, sodium benzoate, sodium citrate dihydrate, sorbitol solution, sucralose

### *Questions or Comments?*

Call 1-888-535-0305 9a.m. – 5p.m. CST

### PRINCIPAL DISPLAY PANEL

NDC 58809-999-01
VanaCof ®

Each 5 mL (1 TEASPOONFUL) CONTAINS:
Chlophedianol Hydrochloride ..... 12.5 mg
Dexchlorpheniramine Maleate ..... 1 mg
Pseudoephedrine Hydrochloride ..... 30mg

NDC 58809-999-01

# VANACOF®

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

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

**Cough Suppressant**
**Antihistamine**
**Nasal Decongestant**

**Sugar Free, Alcohol Free, Dye Free**

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

**THIS BOTTLE IS NOT TO BE DISPENSED TO CONSUMER.**
**Dispense in a tight, light-resistant container with a child-resistant cap.**

N 3 58809 99901 9

US Patent # 9,463,191
US Patent # 9,050,289

**GM Pharmaceuticals, Inc.**

16 fl. oz. (473 mL)

## Drug Facts

| Active ingredients in each 5 mL (1 teaspoonful) | Purpose |
|---|---|
| Chlophedianol Hydrochloride 12.5 mg | Cough Suppressant |
| Dexchlorpheniramine Maleate 1 mg | Antihistamine |
| Pseudoephedrine Hydrochloride 30 mg | Nasal Decongestant |

### Uses

- temporarily relieves these symptoms due to the common cold, hay fever (allergic rhinitis) or other upper respiratory allergies:
  - runny nose
  - sneezing
  - itching of the nose or throat
  - itchy, watery eyes
  - nasal congestion
  - reduces swelling of nasal passages
- cough due to minor throat and bronchial irritation

### 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
- trouble urinating due to an enlarged prostate gland
- glaucoma
- heart disease
- high blood pressure
- thyroid disease
- diabetes

**Ask a doctor before use if you are** taking sedatives or tranquilizers ▶

Distributed by: GM Pharmaceuticals, Inc.
Arlington, TX 76018
201046-01 Rev.0725

PEEL

## Drug Facts (continued)

**When using this product**

- excitability may occur, especially in children
- may cause marked drowsiness
- alcohol, sedatives, and tranquilizers may increase the drowsiness effect
- avoid alcoholic drinks
- use caution 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

- mL= milliliter

| Age | Dose |
|---|---|
| adults and children 12 years of age and over: | 2 teaspoonfuls (10 mL) every 6 hours, not to exceed 8 teaspoonfuls (40 mL) in 24 hours. |
| children 6 to under 12 years of age: | 1 teaspoonful (5 mL) every 6 hours, not to exceed 4 teaspoonfuls (20 mL) in 24 hours. |
| children under 6 years of age: | consult a doctor. |

## Other information

- this packaging is child-resistant.
- store at room temperature of 68°-86°F (20°-30°C) with excursions of 59°-86°F (15°-30°C)

## Inactive ingredients

citric acid anhydrous, flavor, glycerin, propylene glycol, purified water, sodium benzoate, sodium citrate dihydrate, sorbitol solution, sucralose

## Questions?

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

# VANACOF

chlophedianol hydrochloride, dexchlorpheniramine maleate, and pseudoephedrine hydrochloride liquid

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:58809-999 |
| **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 |
| **DEXCHLORPHENIRAMINE MALEATE** (UNII: B10YD955QW) (DEXCHLORPHENIRAMINE - UNII:3Q9Q0B929N) | DEXCHLORPHENIRAMINE 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 |
|---|---|
| **SODIUM BENZOATE** (UNII: OJ245FE5EU) | |
| **ANHYDROUS CITRIC ACID** (UNII: XF417D3PSL) | |
| **GLYCERIN** (UNII: PDC6A3C0OX) | |
| **PROPYLENE GLYCOL** (UNII: 6DC9Q167V3) | |
| **SORBITOL** (UNII: 506T60A25R) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **SODIUM CITRATE** (UNII: 1Q73Q2JULR) | |
| **SUCRALOSE** (UNII: 96K6UQ3ZD4) | |

## 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:58809-999-02 | 12 in 1 TRAY | 04/22/2008 | |
| **1** | | 15 mL in 1 BOTTLE; Type 0: Not a Combination Product | | |
| **2** | NDC:58809-999-01 | 473 mL in 1 BOTTLE; Type 0: Not a Combination Product | 04/22/2008 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | M012 | 04/22/2008 | |

## Labeler - GM Pharmaceuticals, INC (793000860)

Revised: 10/2025 GM Pharmaceuticals, INC