VANACOF DM- dextromethorphan hydrobromide, guaifenesin, and phenylephrine hydrochloride liquid
GM Pharmaceuticals, INC

Vanacof DM

Vanacof DM

NDC 58809-555-08

8 fl. Oz. (240 mL)

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

Dextromethorphan HBr 18 mg

Guaifenesin 200 mg

Phenylephrine HCl 10 mg

Purpose

Cough Suppressant

Expectorant

Nasal Decongestant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive
- temporarily relieves:
  - cough due to minor throat and bronchial irritation as may occur with the common cold or inhaled irritants
  - the intensity of coughing
  - the impulse to cough to help you get to sleep
  - nasal congestion due to a cold

Warnings

Do not use

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

- heart disease

- high blood pressure
- thyroid disease
- diabetes
- a cough that lasts or is chronic such as occurs with smoking, asthma, or emphysema
- a cough that occurs with too much phlegm (mucus)
- difficulty in urination due to enlargement of the prostate gland

When using this product

do not use more than directed

Stop use and ask a doctor if:
- nervousness, dizziness or sleeplessness occurs
- symptoms do not improve within 7 days or are accompanied by a fever, rash or persistent headache. A persistant 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 6 doses in any 24-hour period
- use enclosed dosage cup or tablespoon (TBSP)
- dose as follows or as directed by a doctor

Adults and children 12 years of age and over:	15 mL (1 TBSP) every 4 hours, not to exceed 90 mL (6 TBSP) in a 24 hour period.
Children 6 to under 12 years of age:	7.5 mL (1/2 TBSP) every 4 hours, not to exceed 45 mL (3 TBSP) in a 24 hour period.
Children under 6 years of age:	Consult a doctor.

Other information
- Each 15 mL (TBSP) contains: **Sodium 8 mg.**
- read all product information before using
- store at 20-30°C (68-86°F)

Inactive ingredients

citric acid anhydrous, glycerin, masking agent, propylene glycol, purified water, raspberry flavor, sodium benzoate, sodium citrate dihydrate, sodium saccharin, sorbitol

Questions or Comments?

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

Distributed by:

GM Pharmaceuticals, Inc.

Arlington, TX 76018

KEEP LEAFLET AFTER OPENING

Rev. 09/25

PRINCIPAL DISPLAY PANEL

NDC 58809-555-08

VANACOF DM

Cough

Cold/Congestion

Raspberry Flavor

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Drug Facts

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When using this product do not use more than directed

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occurs
- symptoms do not improve within 7 days or are accompanied by a fever, rash or persistent headache. A persistent cough may be a sign of a serious condition.
- new symptoms occur

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Drug Facts (continued)

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VANACOF DM

dextromethorphan hydrobromide, guaifenesin, and phenylephrine hydrochloride liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:58809-555
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	18 mg in 15 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg in 15 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 15 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SORBITOL (UNII: 506T60A25R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	RASPBERRY	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:58809-555-08	240 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	06/01/2013	

Labeler - GM Pharmaceuticals, INC (793000860)

Revised: 11/2025 GM Pharmaceuticals, INC