

**GNP MUCUS RELIEF PE- guaifenesin/phenylephrine tablet**  
**AmerisourceBergen Drug Corp**

*Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.*

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**DRUG FACTS**

**Active ingredient** - (per tablet)

Guaifenesin 400 mg

Phenylephrine HCl 10 mg

**Purpose**

Guaifenesin.....Expectorant

Phenylephrine HCl.....Nasal decongestant

**Uses**

Temporarily relieves symptoms associated with a cough ,the common cold,hay fever or other upper respiratory allergies.

- helps loosen phlegm (mucus)
- clear nasal passageways
- loosens nasal congestion
- drain bronchial tubes
- shrinks swollen membranes
- clears stuffy nose
- makes coughs more productive
- thin bronchial secretions

**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 are uncertain whether your prescription drug contains an MAOI, ask a health professional.

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

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- excessive phlegm;mucus
- difficulty in urination due to an enlarged prostate gland
- persistent or chronic cough such as occurs with smoking,asthma,chronic bronchitis or emphysema

**Stop use and ask a doctor if**

- nervousness, dizziness or sleeplessness occurs
- symptoms are accompanied by fever, rash, persistent headache or excessive phlegm (mucus)
- cough and congestion do not improve within 7 days or tend to recur.

**These could be signs of a serious condition.**

**If pregnant or breast-feeding**, ask a health professional before use.

**Keep out of the reach of children.** In case of overdose, get medical help or contact a Poison Control Center immediately.

**Directions**

- adults and children 12 years and over: take 1 caplet every 4 hours as needed
- children 6 to under 12 years: take 1/2 caplet every 4 hours as needed
- children under 6 years: consult a doctor

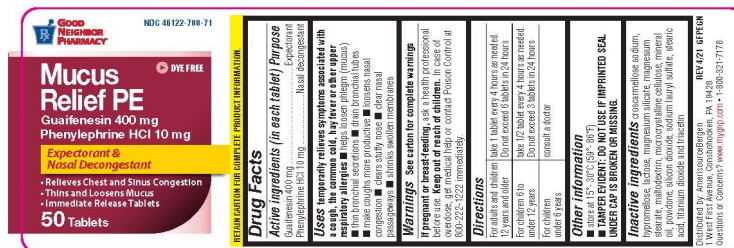
**Do not exceed 6 doses in a 24 hour period or as directed by a doctor.**

**Inactive ingredients**

croscarmellose sodium, Hypromellose, lactose, magnesium silicate, magnesium stearate maltodextrin, microcrystalline cellulose, mineral oil, povidone, silicon dioxide, sodium lauryl sulfate, stearic acid, titanium dioxide, triacetin

**Carton Panel**





## GNP MUCUS RELIEF PE

guaifenesin/phenylephrine tablet

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:46122-700
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>Guaifenesin</b> (UNII: 495W7451VQ) (Guaifenesin - UNII:495W7451VQ)	Guaifenesin	400 mg
<b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>CROSCARMELLOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>LACTOSE</b> (UNII: J2B2A4N98G)	
<b>MALTODEXTRIN</b> (UNII: 7CVR7L4A2D)	

<b>CELLULOSE, MICROCRYSTALLINE</b> (UNII: OP1R32D61U)	
<b>MINERAL OIL</b> (UNII: T5L8T28FGP)	
<b>POVIDONE</b> (UNII: FZ989GH94E)	
<b>MAGNESIUM SILICATE</b> (UNII: 9B9691B2N9)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>SODIUM LAURYL SULFATE</b> (UNII: 368GB5141J)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>TRIACETIN</b> (UNII: XHX3C3X673)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	

### Product Characteristics

<b>Color</b>	white	<b>Score</b>	2 pieces
<b>Shape</b>	OVAL	<b>Size</b>	18mm
<b>Flavor</b>		<b>Imprint Code</b>	PH043
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:46122-700-71	1 in 1 CARTON	01/13/2022	
1		50 in 1 BOTTLE; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	01/13/2022	

**Labeler** - AmerisourceBergen Drug Corp (007914906)

**Registrant** - Reese Pharmaceutical Co (004172052)

### Establishment

Name	Address	ID/FEI	Business Operations
Pharbest		557054835	manufacture(46122-700)

Revised: 7/2023

AmerisourceBergen Drug Corp