## 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



# Mucus Relief PE

DYE FREE

NDC 46122-700-71

Expectorant & Nasal Decongestant

### Drug Facts (continued)

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 Poison Control at 800-222-1222 immediately.

### Directions

For adults and children 12 years and older	take 1 tablet every 4 hours as needed. Do not exceed 6 tablets in 24 hours.	
For children 6 to under 12 years	take 1/2 tablet every 4 hours as needed. Do not exceed 3 tablets in 24 hours.	
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Other information = store at 15°-30°C (59°-86°F)

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SEAL
UNDER CAP IS BROKEN OR MISSING.

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Distributed by:
AmerisourceBergen
1 West First Avenue
Conshohooken, PA 19428
Questions or Concerns?
wewn mygnp. 01-800-321-7178

REV 4/21 GFPEGN

GOOD NEIGHBOR PHARMACY

# DYE FREE Mucus Relief PE

Guaifenesin 400 mg Phenylephrine HCl 10 mg

## **Expectorant &** Nasal Decongestant

- Relieves Chest and Sinus Congestion
- Thins and Loosens Mucus
   Immediate Release
   Tablets

50 Tablets



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   Thins and Loosens Mucus
   Immediate Release Tablets

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### RETAIN CARTON FOR COMPLETE PRODUCT INFORMATION.

## Drug Facts

Active ingredients (in each tablet)

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Expectorant
Nasal decongestant

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## **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	
Guaifenesin (UNII: 495W7451VQ) (Guaifenesin - UNII:495W7451VQ)	Guaifenesin	400 mg	
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg	

Inactive Ingredients		
Ingredient Name	Strength	
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)		
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)		
LACTOSE (UNII: J2B2A4N98G)		
MALTODEXTRIN (UNII: 7CVR7L4A2D)		

CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
MINERAL OIL (UNII: T5L8T28FGP)	
POVIDONE (UNII: FZ 989GH94E)	
MAGNESIUM SILICATE (UNII: 9B9691B2N9)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

Product Characteristics			
Color	white	Score	2 pieces
Shape	OVAL	Size	18mm
Flavor		Imprint Code	PH043
Contains			

P	Packaging				
#	Item Code	Item Code Package Description		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