# MUCUS RELIEF D- guaifenesin and pseudoephedrine hydrochloride tablet, multilayer, extended release H E B

**HEB Mucus Relief D Drug Facts** 

#### Active ingredients (in each extended-release bi-layer tablet)

Guaifenesin 600 mg

Pseudoephedrine HCl 60 mg

#### **Purposes**

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 nasal congestion due to:
- common cold
- hav fever
- upper respiratory allergies
- temporarily restores freer breathing through the nose
- promotes nasal and/or sinus drainage
- temporarily relieves sinus congestion and pressure

## **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
- · trouble urinating due to an enlarged prostate gland
- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema
- cough accompanied by too much phlegm (mucus)

#### When using this product

do not use more than directed

#### Stop use and ask a doctor if

- you get nervous, dizzy, or sleepless
- symptoms do not get better within 7 days, come back or occur with a fever, rash, or persistent headache. These could be signs of a serious illness.

#### 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 (1-800-222-1222).

#### **Directions**

- do not crush, chew, or break tablet
- take with a full glass of water
- this product can be administered without regard for timing of meals
- adults and children 12 years and older: 2 tablets every 12 hours; not more than 4 tablets in 24 hours
- children under 12 years of age: do not use

#### Other information

- · do not use if printed blister unit is broken or torn
- store between 20-25°C (68-77°F)

## Inactive ingredients

carbomer homopolymer type B, colloidal silicon dioxide, FD&C yellow #6 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, sodium starch glycolate

#### Questions or comments?

1-800-719-9260

# Package/Label Principal Display Panel

Compare to Mucinex® D active ingredients

H-E-B<sub>®</sub>

Mucus Relief D

Guaifenesin 600 mg

Pseudoephedrine Hydrochloride 60 mg

Extended-Release Bi-Layer Tablets

Expectorant / Nasal Decongestant

- Clears Nasal/Sinus Congestion
- Thins and Loosens Mucus
- Immediate and Extended Release

12 Hours

**Actual Size** 

18 EXTENDED-RELEASE BI-LAYER TABLETS



#### **MUCUS RELIEF D**

guaifenesin and pseudoephedrine hydrochloride tablet, multilayer, extended release

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59640-621	
Route of Administration	ORAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	600 mg		
PSEUDOEPHEDRINE HYDROCHLORIDE (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F)	PSEUDOEPHEDRINE HYDROCHLORIDE	60 mg		

Inactive Ingredients				
Ingredient Name	Strength			
CARBOMER HOMOPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: HHT01ZNK31)				
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)				
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)				

Product Characteristics					
Color	ORANGE	Score	no score		
Shape	OVAL	Size	17mm		
Flavor		Imprint Code	L6		
Contains					

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59640- 621-89	1 in 1 CARTON	11/30/2023	
1		18 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA214407	11/30/2023	

# **Labeler -** H E B (007924756)

Revised: 1/2024 H E B