MUCUS RELIEF D- guaifenesin, 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 1200 mg

Pseudoephedrine HCl 120 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
- hay 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: 1 tablet every 12 hours; not more than 2 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?

Package/Label Principal Display Panel

Compare to Maximum Strength Mucinex® D active ingredients

H-E-B®

Maximum Strength

Mucus Relief D

Guaifenesin 1200 mg

Pseudoephedrine Hydrochloride 120 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

24 EXTENDED-RELEASE BI-LAYER TABLETS

MUCUS RELIEF D

guaifenesin, pseudoephedrine hydrochloride tablet, multilayer, extended release

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59640-300
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	1200 mg
PSEUDOEPHEDRINE HYDROCHLORIDE (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F)	PSEUDOEPHEDRINE HYDROCHLORIDE	120 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)	
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)	

Product Characteristics

Color	ORANGE	**Score**	no score
Shape	OVAL	**Size**	22mm
Flavor		**Imprint Code**	L12
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59640-300-62	4 in 1 CARTON	11/30/2023	
1		6 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)