MUCUS D- guaifenesin and pseudoephedrine hydrochloride tablet, multilayer, extended release Meijer, Inc.

Meijer Distribution, Inc. Mucus-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
- 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: 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?

1-800-719-9260

Package/Label Principal Display Panel

 $\text{meijer}_{\mathbb{R}}$

Compare to Maximum Strength Mucinex[®] D active ingredients

MAXIMUM STRENGTH

mucus-D

Guaifenesin 1200 mg & Pseudoephedrine Hydrochloride 120 mg Extended-Release Bi-Layer Tablets

Expectorant | Nasal Decongestant

12 HOUR | Clears Nasal/Sinus Congestion

Thins And Loosens Mucus

Immediate and Extended-Release

24 Extended-Release Bi-Layer Tablets | actual size



MUCUS D

guaifenesin and pseudoephedrine hydrochloride tablet, multilayer, extended release

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79481-0119
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				

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:79481- 0119-0	4 in 1 CARTON	08/22/2023		
1		3 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	08/22/2023	

Labeler - Meijer, Inc. (006959555)

Revised: 9/2023 Meijer, Inc.