# MUCUS RELIEF D- guaifenesin and pseudoephedrine hcl tablet, extended release

Wal-mart Stores, Inc.

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#### **Drug Facts**

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

Guaifenesin USP, 1200
mg......Expectorant
Pseudoephedrine HCl USP, 120 mg.....Nasal
Decongestant

#### **Purpose**

**Expectorant and 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:
  - o 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 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

- tamper evident: do not use if carton is open or if printed seal on blister is broken or missing
- store between 20° 25°C (68° 77°F)

# **Inactive ingredients**

FD & C yellow #6, hypromellose, lactose monohydrate, magnesium stearate, microcrystalline cellulose, povidone, silicon dioxide.

#### **Questions?**

call **1-888-287-1915** 

You may also report side effect to this phone number.

## **Principal Display Panel**



#### **MUCUS RELIEF D**

guaifenesin and pseudoephedrine hcl tablet, extended release

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-261(NDC:55111-799)
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
Guaifenesin (UNII: 495W7451VQ) (Guaifenesin - UNII:495W7451VQ)	Guaifenesin	1200 mg	
<b>Pseudoephedrine Hydrochloride</b> (UNII: 6V9V2RYJ8N) (Pseudoephedrine - UNII:7CUC9DDI9F)	Ps eudoephedrine Hydrochloride	120 mg	

Inactive Ingredients				
Ingredient Name	Strength			
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)				
HYPROMELLOSES (UNII: 3NXW29V3WO)				
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)				
POVIDONE (UNII: FZ989GH94E)				
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				

Product Characteristics			
Color	ORANGE	Score	no score
Shape	CAPSULE	Size	22mm
Flavor		Imprint Code	RDY;799
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:49035- 261-24	4 in 1 CARTON	04/03/2020	04/03/2020	
1		6 in 1 BLISTER PACK; Type 0: Not a Combination Product			
2	NDC:49035- 261-12	2 in 1 CARTON	04/03/2020		
2		12 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	ANDA208369	04/03/2020	

# Labeler - Wal-mart Stores, Inc. (051957769)

Revised: 5/2023 Wal-mart Stores, Inc.