EQUATE MUCUS RELIEF- guaifenesin tablet, multilayer, extended release
WALMART INC.

Wal-Mart Mucus Relief Drug Facts

Active ingredient (in each extended-release tablet)

Guaifenesin 1200 mg

Purpose

Expectorant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive

Warnings

Do not use

- for children under 12 years of age

Ask a doctor before use if you have

- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema
- cough accompanied by too much phlegm (mucus)

Stop use and ask a doctor if

- cough lasts more than 7 days, comes back, or occurs with fever, rash, or persistent headache. These could be signs of a serious illness.

If pregnant or breast-feeding,

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 extended-release tablet
- take with a full glass of water

- this product can be administered without regard for timing of meals
- adults and children 12 years of age and over: 1 extended-release tablet every 12 hours. Do not exceed 2 extended-release tablets in 24 hours.
- children under 12 years of age: do not use

Other information

- store at 20-25°C (68-77°F)

Inactive ingredients

carbomer homopolymer type B, copovidone, FD&C blue #1 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, silicon dioxide, sodium starch glycolate

Questions or comments?

1-888-287-1915

Package/Label Principal Display Panel

equate™

Compare Maximum Strength Mucinex® active ingredient

Mucus Relief

Guaifenesin Extended-Release Tablets, 1200 mg

Expectorant

MAXIMUM STRENGTH

• Relieves chest congestion

• Thins and loosens mucus

• Immediate and extended release

12 HOUR

1200 mg

14 EXTENDED-RELEASE TABLETS

Actual Size

EQUATE MUCUS RELIEF

guaifenesin tablet, multilayer, extended release

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79903-327
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	1200 mg

Inactive Ingredients	
Ingredient Name	**Strength**
CARBOMER HOMOPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: HHT01ZNK31)	
COPOVIDONE K25-31 (UNII: D9C330MD8B)	
FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)	

Product Characteristics

Color	BLUE	**Score**	no score
Shape	OVAL	**Size**	22mm
Flavor		**Imprint Code**	L4S1
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79903-327-14	2 in 1 CARTON	03/31/2025	
1		7 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:79903-327-28	4 in 1 CARTON	03/31/2025	
2		7 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	ANDA078912	03/31/2025	

Labeler - WALMART INC. (051957769)

Revised: 4/2025 WALMART INC.