

MUCUS RELIEF DM- dextromethorphan hydrobromide/guaifenesin tablet RITE AID

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

280R Rite aid 11822 7894 Mucus Relief DM

DRUG FACTS

Active ingredients (in each immediate-release tablet)

Dextromethorphan HBr 20 mg

Guaifenesin 400 mg

Purpose

Cough suppressant

Expectorant

Uses

- helps loosen phlegm (mucus)
- helps thin bronchial secretions to make coughs more productive
- temporarily relieves cough due to minor throat and bronchial irritation associated with the common cold

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

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

When using the product

- **do not exceed recommended dosage**
- do not use for more than 7 days

Stop use and ask a doctor if

- cough lasts for more than 7 days, recurs, or is accompanied by fever, rash, or persistent headache. These could be signs of a serious condition.

If pregnant or breast-feeding, ask a healthcare professional before use

Keep out of reach of children. In case of overdose, get medical help or contact a

Poison Control Center right away.

Directions

- adults and children 12 years of age and older: take 1 tablet every 4 hours with a full glass of water while symptoms persist. Do not exceed 6 doses in 24 hours.
- children under 12 years of age: do not use

Other information

- store at 25°C (77°F) excursions permitted between 15°-30°C (59°-86°F)
- keep in a dry place and do not expose to excessive heat

Inactive ingredients colloidal silicon dioxide, magnesium stearate, maltodextrin, microcrystalline cellulose, povidone, silicon dioxide, sodium starch glycolate, stearic acid

Questions or comments?

Call 1-877-290-4008



NDC 11822-7894-3

MUCUS RELIEF DM

**GUAIFENESIN 400 mg
DEXTROMETHORPHAN HBr 20 mg**

**EXPECTORANT
& COUGH SUPPRESSANT**

Controls cough
Thins & loosens mucus
Immediate-release

30

IMMEDIATE-RELEASE TABLETS

READ AND KEEP CARTON FOR COMPLETE WARNINGS AND INFORMATION

Uses

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- temporarily relieves cough due to minor throat and bronchial irritation associated with the common cold

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Directions

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- children under 12 years of age: do not use

Other information

- **TAMPER EVIDENT: DO NOT USE THIS PRODUCT IF THE IMPRINTED FOIL SEAL OVER THE MOUTH OF THE BOTTLE IS CUT, TORN, BROKEN OR MISSING.**
- store at 25°C (77°F) excursions permitted between 15°-30°C (59°-86°F)
- keep in a dry place and do not expose to excessive heat

Questions or comments? Call 1-877-290-4008

DISTRIBUTED BY: SATISFACTION GUARANTEE:
RITE AID
30 HUNTER LANE
CAMP HILL, PA 17011
www.riteaid.com

Lot No.: _____
Exp. Date: _____

280R 0821

MUCUS RELIEF DM

dextromethorphan hydrobromide/guaifenesin tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-7894
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg

Inactive Ingredients

Ingredient Name	Strength
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POVIDONE (UNII: FZ989GH94E)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics

Color	white	Score	no score
Shape	OVAL (Oval shaped caplet)	Size	16mm
Flavor		Imprint Code	TCL280
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-7894-3	30 in 1 CARTON; Type 0: Not a Combination Product	01/01/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	01/01/2022	

Labeler - RITE AID (014578892)

Registrant - TIME CAP LABORATORIES, INC. (037052099)

Establishment

Name	Address	ID/FEI	Business Operations
TIME CAP LABORATORIES, INC.		037052099	manufacture(11822-7894)

Revised: 12/2021

RITE AID