

ACTIDOM DMX- dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride solution
ACTIPHARMA, LLC

ACTIDOM DMX

Drug Facts

Active ingredients (in each 5 mL tsp)

Dextromethorphan HBr, 30 mg
Guaifenesin, 200 mg
Phenylephrine HCl, 10 mg

Purposes

Cough suppressant
Expectorant
Nasal decongestant

Uses

- Helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes
- Temporarily relieves these symptoms occurring with a cold:
- nasal congestion • cough due to minor throat and bronchial irritation.

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 • cough that occurs with too much phlegm (mucus) • cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis or emphysema.

Ask a doctor or pharmacist before use if you are taking any other oral nasal decongestant or stimulant.

When using this product • do not exceed recommended dosage

Stop use and ask a doctor if

- you get nervous, dizzy or sleepless
- symptoms do not get better within 7 days or are accompanied by fever
- coughs lasts more than 7 days, comes back, or is accompanied by fever, rash, or a

persistent headache. These could be signs of a serious condition.

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.

Directions

- do not take more than 4 doses in any 24-hour period.

AGE	DOSE
Adults and Children 12 years of age and over	Take 1 teaspoonfuls (5 mL) every 6 hours
Children under 12 years of age	ask a doctor

Other information

- Tamper Evident Feature: Do not use if safety seal is torn, cut or opened.
- Store at controlled room temperature 15° - 30°C (59° - 86°F).
- Avoid excessive heat or humidity.

Inactive ingredients

Citric acid, D&C Red #40, FD&C Blue #1, flavor, glycerin, methylparaben, propylene glycol, propylparaben, purified water, sodium citrate, sucralose.

Questions or comments?

call weekdays from 8AM to 4PM AST at **1.787.608.0882**

Contains the same active ingredients as Dometuss®-DMX*

SUGAR FREE

ALCOHOL FREE

Grape Flavor

Manufactured in USA with imported ingredients for ActiPharma. San Juan, PR 00917.
www.actipharma.net.

* Dometuss® DMX is a registered trademark of Domel Laboratories. This product is not manufactured, distributed or marketed by Domel Laboratories.

Packaging

NDC 63102-110-16

ACTIDOM[®] DMX

Contains the same active ingredients as Dometuss[®] DMX*

**COUGH SUPPRESSANT
EXPECTORANT
NASAL DECONGESTANT**

**SUGAR FREE
ALCOHOL FREE**

Grape Flavor



16 Fl.oz. (473 mL)

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 Guaifenesin, 200 mg.....Expectorant
 Phenylephrine HCl, 10 mg..... Nasal decongestant

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Drug Facts (continued)

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ACTIDOM DMX

dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63102-110
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	30 mg in 5 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg in 5 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	

SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)

SUCRALOSE (UNII: 96K6UQ3ZD4)

Product Characteristics

Color	purple (Clear)	Score	
Shape		Size	
Flavor	grape	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63102-110-16	474 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/13/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	08/13/2015	

Labeler - ACTIPHARMA, LLC (079340948)

Revised: 6/2024

ACTIPHARMA, LLC