M-END DMX- dexbrompheniramine maleate, dextromethorphan hydrobromide, pseudoephedrine hydrochloride liquid
R. A. McNeil Company

M-END DMX

Drug Facts

Active ingredients
(in each 5 mL teaspoonful)
Dexbrompheniramine Maleate 0.667 mg
Dextromethorphan HBr 10 mg
Pseudeophedrine HCl 20 mg

Purpose

Antihistamine

Antitussive

Cough Suppressant

Nasal Decongestant

Uses

temporarily relieves these symptoms due to the common cold, hay fever (allergic rhinitis) or other upper respiratory allergies:

- runny nose
- sneezing
- itching of the nose or throat
- itchy, watery eyes
- cough due to minor throat and bronchial irritation
- nasal congestion
- reduces swelling of nasal passages

Warnings

Do not exceed recommended dosage.

Do not use this product

- 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

- a breathing problem such as emphysema or chronic bronchitis
- glaucoma
- trouble urinating due to an enlarged prostate gland
- a cough that lasts or is chronic such as occurs with smoking, asthma or emphysema
- a cough that occurs with too much phlegm (mucus)
- heart disease
- high blood pressure
- thyroid disease
- diabetes

Ask a doctor or pharmacist before use if you are taking sedatives or tranquilizers.

When using this product
- excitability may occur, especially in children
- may cause marked drowsiness
- avoid alcoholic drinks
- alcohol, sedatives, and tranquilizes may increase the drowsiness effect
- be careful when driving a motor vehicle or operating machinery

Stop use and ask a doctor if
- cough or nasal congestion persists for more than 1 week, tends to recur, or is accompanied by a fever, rash or persistent headache. A persistent cough may be a sign of a serious condition.
- nervousness, dizziness, or sleeplessness occur
- new symptoms occur

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 exceed 4 doses in a 24 hour period.

Adults and children 12 years of age and over:	3 teaspoonfuls (15 mL) every 6 hours, not to exceed 12 teaspoonfuls in 24 hours
Children 6 to under 12 years of age:	1 1/2 teaspoonfuls (7.5 mL) every 6 hours, not to exceed 6

	teaspoonfuls in 24 hours
Children under 6 years of age:	Consult a Physician.

Other information

Store at 59°-86°F (15°-30°C)

Inactive ingredients

Bitter Mask, Citric Acid, Glycerin, Propylene Glycol, Purified Water, Sodium Citrate, Sodium Saccharin, Sorbitol, Sucralose, Tutti Frutti Flavor.

Questions? Comments?

Call 1-423-493-9170

Product Packaging

Tamper evident by foil seal under cap. Do not use if foil seal is broken or missing.

THIS BOTTLE IS NOT TO BE DISPENSED TO THE CONSUMER.

Dispense in a tight, light-resistant container with a child resistant cap.

Manufactured for:
R.A. McNeil Company
1150 Latta Street
Chattanooga, TN 37406-3738

Rev. 12/14

NDC 12830-0816-16

M-END DMX

Antihistamine • Antitussive • Nasal Decongestant

Alcohol Free • Gluten Free Sugar Free • Dye Free

EACH 5 mL (1 TEASPOONFUL) CONTAINS:

Dexbrompheniramine Maleate	0.667 mg
Dextromethorphan HBr	10 mg
Pseudoephedrine HCl	20 mg

Tutti-Frutti Flavor

ONE PINT (473 mL)

Mfg. for R.A. McNeil Company
Chattanooga, TN 37406-3738

Rev. 12/14

Drug Facts

Lift Here

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	Nasal Decongestant

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M-END DMX

dexbrompheniramine maleate, dextromethorphan hydrobromide, pseudoephedrine hydrochloride liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:12830-816
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXBROMPHENIRAMINE MALEATE (UNII: BPA9UT29BS) (DEXBROMPHENIRAMINE - UNII:75T64B71RP)	DEXBROMPHENIRAMINE MALEATE	0.667 mg in 5 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 5 mL
PSEUDOEPHEDRINE HYDROCHLORIDE (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F)	PSEUDOEPHEDRINE HYDROCHLORIDE	20 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	TUTTI FRUTTI	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:12830-816-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/13/2011	
2	NDC:12830-816-30	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/13/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	10/13/2011	

Labeler - R. A. McNeil Company (008305220)

Revised: 10/2025 R. A. McNeil Company