ABATUSS DMX- dexchlorpheniramine maleate, dextromethorphan hydrobromide, pseudoephedrine hydrochloride liquid
Kramer Novis

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.

ABATUSS DMX Antihistamine Cough Suppressant Nasal Decongestant GRAPE Flavor

Drug Facts

Active ingredients (per 5 mL)

Dexchlorpheniramine Maleate 1 mg
Dextromethorphan HBr 15 mg
Pseudoephedrine HCl 30 mg

Purpose

Antihistamine
Cough Suppressant
Nasal Decongestant

Uses

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

- sneezing
- itchy nose or throat
- runny nose
- itchy, watery eyes
- nasal congestion
- temporarily controls cough due to minor throat and bronchial irritation associated with inhaled irritants
- temporarily restores freer breathing through nose

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 • a breathing problem such as emphysema or chronic bronchitis • glaucoma • difficulty in urination due to enlarged prostate gland • persistent or chronic cough such as occurs with smoking, asthma, or emphysema • cough accompanied by excessive phlegm (mucus)

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

When using this product • do not exceed recommended dosage • may cause drowsiness • alcohol, sedatives, and tranquilizers may increase the drowsiness effect • avoid alcoholic beverages • use

caution when driving a motor vehicle or operating machinery marked may occur • excitability may occur especially in children

Stop use and ask a doctor if • nervousness, dizziness, or sleeplessness occur • cough persists for more than 1 week, tends to recur, or is accompanied by fever, rash, or persistent headache. A persistent cough may be a sign of a serious condition. • symptoms do not improve within 7 days or are accompanied by fever

If pregnant or breastfeeding 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:

Adults and children 12 years of age and older	2 teaspoonfuls (10 mL) every 6 hours, not to exceed 8 teaspoonfuls in 24 hours or as directed by a doctor
Children 6 to under 12 years of age	1 teaspoonful (5 mL) every 6 hours, not to exceed 4 teaspoonfuls in 24 hours or as directed by a doctor
Children 2 to under 6 years of age	Consult a doctor

Other information

- Store at controlled room temperature 15°C-30°C (59°F-86°F)
- Tamper evident by foil seal under cap. Do not use if foil seal is broken or missing

Inactive ingredients:

Citric acid, FD&C blue #1, FD&C Red #40, grape flavor, methylparaben, monoammonium glycyrrhizinate, potassium citrate, potassium sorbate, propylene glycol, propylparaben, purified water, sorbitol, sucralose.

*Contains the same active ingredients as Deltuss DMX®

Antihistamine
Cough Suppressant
Nasal Decongestant

Each 5 mL contains:

Dexchlorpheniramine Maleate .....1 mg
Dextromethorphan HBr.....15 mg
Pseudoephedrine HCl .....30 mg

• Alcohol Free • Sugar Free

• Gluten Free

Grape Flavor

Manufactured in the USA for Kramer Novis, San Juan, PR 00917. T: (787) 767-2072 www.kramernovis.com

* Deltuss DMX® is a registered trademark of Deliz Pharmaceutical Corp. This product is not manufactured, distributed or marketed by Deliz Pharmaceutical Corp.

Packaging

ABATUSS DMX

dexchlorpheniramine maleate, dextromethorphan hydrobromide, pseudoephedrine hydrochloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52083-625
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DEXCHLORPHENIRAMINE MALEATE (UNII: B10YD955QW) (DEXCHLORPHENIRAMINE - UNII:3Q9Q0B929N)	DEXCHLORPHENIRAMINE MALEATE	1 mg in 5 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	15 mg in 5 mL
PSEUDOEPHEDRINE HYDROCHLORIDE (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F)	PSEUDOEPHEDRINE HYDROCHLORIDE	30 mg in 5 mL

Inactive Ingredients	
Ingredient Name	**Strength**
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	

METHYLPARABEN (UNII: A2I8C7HI9T)	
AMMONIUM GLYCYRRHIZATE (UNII: 3VRD35U26C)	
POTASSIUM CITRATE (UNII: EE90ONI6FF)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	GRAPE (Grape Flavor)	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52083-625-16	1 in 1 CARTON	05/25/2014	
1		473 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	05/25/2014	

Labeler - Kramer Novis (090158395)

Revised: 11/2018 Kramer Novis