CAPRON DM- dextromethorphan hbr and pyrilamine maleate liquid Capital Pharmaceutical, LLC

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.

Capron DM Liquid

Drug Facts

Active Ingredients

(in each 5 mL teaspoonful)

Dextromethorphan HBr 7.5 mg

Purpose

Antitussive

Active Ingredients

(in each 5 mL teaspoonful)

Pyrilamine Maleate 7.5 mg

Purpose

Antihistamine

Uses

temporarily relieves these symptoms due to 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 associated with a cold
- alleviates cough to help you sleep
- nonnarcotic cough suppressant for the relief of cough

Warnings

Do not exceed recommended dosage.

 a persistent cough may be a sign of a serious condition. If cough persist for more than 1 week, tends to recur, or is accompanied by a fever, rash, or persistent headache, consult a doctor.

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
- for persistent or chronic cough such as occurs with smoking, asthma, or emphysema, or if cough is accompanied by excessive phlegm (mucus) unless directed by a doctor

Ask a doctor before use if you have

- a breathing problem such as emphysema or chronic bronchitis
- glaucoma
- difficulty in urination due to enlargement of the prostate gland

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

When using this product

- excitability may occur, especially in children
- may cause marked drowsiness
- sedatives and tranquilizers may increase drowsiness effect
- avoid alcoholic beverages
- use caution when driving a motor vehicle or operating machinery

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of accidental overdose, seek professional help or contact a Poison Control Center immediately.

Directions

Do not exceed recommended dosage.

Adults and children 12 years of age and over:	4 teaspoonfuls every 6-8 hours, not to exceed 16 teaspoonfuls in 24 hours, or as directed by a doctor
Children 6 to under 12 years of age:	2 teaspoonfuls every 6-8 hours, not to exceed 8 teaspoonfuls in 24 hours, or as directed by a doctor
Children 2 to under 6 years of age	1 teaspoonful every 6-8 hours, not to

Children under 2 years of age:

Consult a doctor

Inactive ingredients

Citric Acid, Flavor, Magnasweet, Methyl Paraben, Potassium Citrate, Potassium Sorbate, Propyl Paraben, Propylene Glycol, Purified Water, Sorbitol, Sucralose

Questions or Comments?

Call 614.638.4622

PRINCIPAL DISPLAY PANEL



NDC 29978-127-16

Capron DM

Liquid

Candy Apple Flavor

16 oz. (473 mL)

PRINCIPAL DISPLAY PANEL

NDC 29978-127-15 Capron DM Liquid Candy Apple Flavor 1 5 mL



CAPRON DM

dextromethorphan hbr and pyrilamine maleate liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:29978-127
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	7.5 mg in 5 mL	
PYRILAMINE MALEATE (UNII: R35D29L3ZA) (PYRILAMINE - UNII:HPE317O9TL)	PYRILAMINE MALEATE	7.5 mg in 5 mL	

Inactive Ingredients		
Ingredient Name	Strength	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		
AMMONIUM GLYCYRRHIZATE (UNII: 3VRD35U26C)		
METHYLPARABEN (UNII: A2I8C7HI9T)		
POTASSIUM CITRATE (UNII: EE90ONI6FF)		

POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics			
Color		Score	
Shape		Size	
Flavor	APPLE (Candy Apple)	Imprint Code	
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:29978-127- 15	6 in 1 BOX	06/20/2013	
1		15 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:29978-127- 16	1 in 1 BOX	06/20/2013	
2		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	06/20/2013	

Labeler - Capital Pharmaceutical, LLC (831545541)

Revised: 1/2023 Capital Pharmaceutical, LLC