

SUPRESS A- dextbromopheniramine maleate, dextromethorphan hbr, phenylephrine hcl syrup
Dextrum Laboratories, Inc

SUPRESS A

Active Ingredient Section

Active Ingredient Section

DEXBROMPHENIRAMINE MALEATE; DEXTROMETHORPHAN HYDROBROMIDE;
PHENYLEPHRINE HYDROCHLORIDE



Inactive Ingredient Section

Inactive Ingredient Section:

citric Acid, flavor, glycerin, methylparaben, polysorbate, propylene glycol, propylparaben, purified water, sodium citrate, sucralose.



Do not use section

Do not use section:

Do not use more than 6 doses in any 24-hour period, repeat every 4 hour, measure with the dosage device provided.

Do not use with any other device, children 6 to under 12 years of age L 1 ml, Children under 6 year of age, consult a doctor.



Keep out of reach of children section

Keep out of reach of children section:

In case of accidental overdose, get medical help or contact a Poison Control Center right away.

NDC 52083-057-01

SUPRESS[®]

- Antihistamine
- Cough Suppressant
- Nasal Decongestant

PEDIATRIC DROPS

Sugar, Dye & Alcohol Free

Net Content: 30 mL (1 Fl. Oz.)

Drug Facts

Active Ingredients (in each 1 mL):

- Antihistamine: **Dechlorpheniramine Maleate, 1 mg.**
- Cough Suppressant: **Dextromethorphan HBr, 10 mg.**
- Nasal Decongestant: **Phenylephrine HCl, 5 mg.**

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 related irritants.

Temporarily restores free breathing through nose.

Warnings: Keep out of reach of children. In case of accidental overdose, get medical help or contact a Poison Control Center right away.

Directions: Do not use more than 6 doses in any 24-hour period. Repeat every 4 hours. Do not use with the cough suppressant. Do not use with any other drugs. Children 6 to under 12 years of age: 1 mL. Children under 6 years of age, consult a doctor.

Other information: Tamper Evident Feature: Do not use if cap seal is torn, broken or missing. Store at controlled room temperature 15-30°C (59-86°F). Avoid exposure to heat or humidity.

Inactive Ingredients: citric acid, fennel, glycerin, methoxyphenol, polyethylene glycol, propylene glycol, propylparaben, purified water, sodium citrate, sucrose.

Questions or comments? Call weekdays from 8 am to 4 pm AST at 1.888.762.2022

Manufactured in the USA for Kramer Novels, San Juan, PR 00917. www.kramernovels.com

RETAIN CARTON FOR FULL PRESCRIBING INFORMATION **Rev. 10/22**

Purpose Section

Purpose Section:

Antihistamine, Cough suppressant, Nasal descongellant



Dosage & Administration Section

Dosage & Administration Section:

Do not use more than 6 doses in any 24-hour period, repeat every 4 hour, measure with the dosage device provided.

Do not use with any other device

children 6 to under 12 years of age 1 ml, Children under 6 year of age, consult a doctor.



Indication and usage Section

Indication and usage Section:

Use 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.



Warning Section

Warning Section:

Keep out of reach of children. In case of overdose,, get medical help or contact Poison Control Center right away.



Question Section

Question Section:

Call weekdays from 8 am to 4 pm AST at 1.787.767.2072



Package Label. Principal Display Panel

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NDC 52083--057-01

SUPRESS A

- Antihistamine
- Cough Suppressant
- Nasal Decongestant

PEDIATRIC DROPS

Sugar, Dye and Alcohol Free

PSEUDOEPHEDRINE & PPA FREE

Net content: 30ml (1Fl, Oz)

Kramer Novis



SUPRESS A

dexbromopheniramine maleate, dextromethorphan hbr, phenylephrine hcl syrup

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65852-020
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXBROMPHENIRAMINE MALEATE (UNII: BPA9UT29BS) (DEXBROMPHENIRAMINE - UNII: 75T64B71RP)	DEXBROMPHENIRAMINE MALEATE	1 mg in 1 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1W5297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 1 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII: 7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLPARABEN (UNII: A2I8C7HI9T)	

CITRIC ACID (UNII: 2968PHW8QP)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	GRAPE	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65852-020-01	1 in 1 CARTON	08/01/2012	
1		30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	08/01/2012	

Labeler - Dextrum Laboratories, Inc (007392322)

Establishment

Name	Address	ID/FEI	Business Operations
Dextrum Laboratories, Inc		007392322	manufacture(65852-020)

Revised: 10/2025

Dextrum Laboratories, Inc