COLD AND COUGH RELIEF CHILDREN- brompheniramine maleate, dextromethorphan hbr, phenylephrine hcl liquid Strategic Sourcing Services LLC

Drug Facts

Active ingredients (in each 10 mL)

Brompheniramine maleate 2 mg

Dextromethorphan HBr 10 mg

Phenylephrine HCl 5 mg

Purposes

Antihistamine

Cough suppressant

Nasal decongestant

Uses

- temporarily relieves cough due to minor throat and bronchial irritation occurring with a cold, and nasal congestion due to the common cold, hay fever or other upper respiratory allergies
- temporarily relieves these symptoms due to hay fever (allergic rhinitis):
 - runny nose
 - sneezing
 - itchy, watery eyes
 - itching of the nose or throat
- temporarily restores freer breathing through the nose

Warnings

Do not use

- to sedate a child or to make a child sleepy
- 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

- glaucoma
- trouble urinating due to an enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- a breathing problem or persistent or chronic cough that lasts such 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
- taking sedatives or tranquilizers

When using this product

- do not use more than directed
- marked drowsiness may occur
- avoid alcoholic beverages
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

Stop use and ask a doctor if

- nervousness, dizziness or sleeplessness occurs
- symptoms do not improve within 7 days or are accompanied by fever
- cough lasts more than 7 days, comes back, or is accompanied by fever, rash, or 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 (1-800-222-1222) right away.

Directions

- do not take more than 6 doses in a 24-hour period
- measure only with dosing cup provided. Do not use any other dosing device
- keep dosing cup with product
- mL = milliliter

age	dose
adults and children 12 years and over	20 mL every 4 hours
children 6 to 11 years	10 mL every 4 hours
children under 6 years	do not use

Other information

• each 10 mL contains: sodium 5 mg

• store between 20-25°C (68-77°F). Do not refrigerate

Inactive ingredients

citric acid, FD&C blue #1, FD&C red 40, flavor, glycerin, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol, sucralose

Questions or comments?

Call **1-877-753-3935** Monday-Friday 9AM-5PM EST

Principal Display Panel

COMPARE TO THE ACTIVE INGREDIENTS IN CHILDREN'S DIMETAPP® COLD & COUGH Children's

Cold & Cough Relief

Relieves itchy, watery eyes, cough, runny nose, sneezing, stuffy nose, itching of the nose or throat

For ages 6 years and over

Alcohol-free

BROMPHENIRAMINE MALEATE 2 MG

ANTIHISTAMINE

DEXTROMETHORPHAN HBr 10 MG

COUGH SUPPRESSANT

PHENYLEPHRINE HCI 5 MG

NASAL DECONGESTANT

GRAPE FLAVOR

fl oz (mL)

Dosing Cup Included

*This product is not manufactured or distributed by Pfizer Consumer Healthcare, distributor of Children's Dimetapp® Cold & Cough.

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPENED OR IF PRINTED SAFETY SEAL AROUND BOTTLE OR UNDER CAP IS BROKEN OR MISSING.

Distributed by McKesson Corp.

6555 State Highway 161 Irving, TX 75039

www.sunmarkbrand.com

Package Label



SUNMARK Children's Cold & Cough Grape Flavor

COLD AND COUGH RELIEF CHILDREN brompheniramine maleate, dextromethorphan hbr, phenylephrine hcl liquid **Product Information Product Type** HUMAN OTC DRUG Item Code (Source) NDC:70677-0121 **Route of Administration** ORAL **Active Ingredient/Active Moiety Ingredient Name Basis of Strength** Strength BROMPHENIRAMINE MALEATE (UNII: IXA7C9ZN03) (BROMPHENIRAMINE -BROMPHENIRAMINE 2 ma

UNII:H57G17P2FN)	MALEATE	in 10 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 10 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 10 mL

Inactive Ingredients	
Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZ B9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics		
Color		Score
Shape		Size
Flavor	GRAPE	Imprint Code
Contains		

Packaging

# Item Cod	Package Description	Marketing Start Date	Marketing End Date
1 NDC:70677- 0121-1	1 in 1 BOX	10/30/2020	
1	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
Marketin	g Information		
Marketin Marketin Category	g Application Number or Monograp	oh Marketing Start Date	Marketing End Date

Labeler - Strategic Sourcing Services LLC (116956644)

Revised: 5/2024