

**TARGET CHILDRENS COLD AND COUGH- brompheniramine maleate,
dextromethorphan hbr, phenylephrine hcl liquid
TARGET CORPORATION**

TARGET Children's Cold & Cough 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:
- Nasal congestion
 - runny nose
 - cough
 - sneezing
 - itchy, watery eyes due to hay fever
 - 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
- trouble urinating due to an enlarged prostate gland

- glaucoma
- cough that occurs with too much phlegm (mucus)
- a breathing problem or persistent or chronic cough that lasts such as 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**
- may cause marked drowsiness
- 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 get better 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 right away at 1-800-222-1222.

Directions

- do not take more than 6 doses in any 24-hour period
- measure only with dosage cup provided
- keep dosage cup with product
- ml = milliliter

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

Other information

- **each 10 ml contains:** sodium 6 mg
- store at room temperature.

Inactive ingredients

citric acid, FD&C blue no. 1, FD&C red no. 40, artificial flavor, glycerin, propylene glycol, propyl gallate, purified water, sodium benzoate, sodium citrate, sorbitol solution, sucralose

Questions or comments?

1-866-467-2748

PRINCIPAL DISPLAY PANEL

Compare to the active ingredients in Children's Dimetapp® Cold & Cough*

NDC 82442-645-04

Children's

Cold & Cough

Brompheniramine maleate(Antihistamine)

Dextromethorphan HBr(Cough Suppressant)

Phenylephrine HCl(Nasal Decongestant)

Relieves + comforts

- **Stuffy, runny nose**
- **Itchy, watery Eyes**
- **Sneezing**
- **Cough**

For Ages 6 Years and Over

No Added Alcohol

Grape Flavor

Naturally and Artificially Flavored

4 FL OZ (118 ml)

Distributed by:

This product is not manufactured or distributed by Foundation Consumer Brands, LLC, the distributor of Children's Dimetapp® Cold & Cough.

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BROMPHENIRAMINE MALEATE (UNII: IXA7C9ZN03) (BROMPHENIRAMINE - UNII:H57G17P2FN)	BROMPHENIRAMINE MALEATE	2 mg 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: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYL GALLATE (UNII: 8D4SNN7V92)	
WATER (UNII: 059QF0K00R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics

Color	purple (purple liquid)	Score	
Shape		Size	
Flavor	GRAPE (grape flavor and odor)	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82442-645-04	1 in 1 CARTON	06/17/2024	
1		118 mL in 1 BOTTLE; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	06/17/2024	

Revised: 7/2024

TARGET CORPORATION