

TARGET CHILDRENS COUGH RELIEF- dextromethorphan hydrobromide and guaifenesin liquid
TARGET CORPORATION

Target Children's Cough Relief Cherry Flavor 4FL OZ

Drug Facts

<i>Active ingredients (in each 5 mL)</i>	<i>Purpose</i>
Dextromethorphan HBr 5 mg	Cough suppressant
Guaifenesin 100 mg	Expectorant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive
- temporarily relieves:
 - cough due to minor throat and bronchial irritation as may occur with the common cold or inhaled irritants
 - the intensity of coughing
 - the impulse to cough to help your child get to sleep

Warnings

Do not use in a child who is 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 child's prescription drug contains an MAOI, ask a doctor or pharmacist before giving this product.

Ask a doctor before use if the child has

- cough that occurs with too much phlegm (mucus)
- persistent or chronic cough such as occurs with asthma

When Using

this product do not use more than directed

Stop use and ask a doctor if cough lasts more than 7 days, comes back, or occurs with fever, rash, or persistent headache. These could be signs of a serious condition.

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 give more than 6 doses in any 24-hour period
- measure only with dosing cup provided
- do not use dosing cup with other products
- dose as follows or as directed by a doctor

Age	Dose
children 6 years to under 12 years	5 mL - 10 mL every 4 hours
children 4 years to under 6 years	2.5 mL - 5 mL every 4 hours
children under 4 years	do not use

Other information

- **each 5 mL contains:** sodium 2 mg
- store at room temperature
- do not refrigerate

Inactive ingredients

anhydrous citric acid, edetate disodium, FD&C red #40, flavors, potassium citrate, propylene glycol, purified water, sodium benzoate, sorbitol, sucralose, xanthan gum

Questions or comments?

1-866-467-2748

TAMPER-EVIDENT: DO NOT USE IF PRINTED INNER SEAL UNDER CAP IS TORN OR MISSING

PRINCIPAL DISPLAY PANEL - 118 mL Bottle Carton

Compare to the Active Ingredients in Children's Mucinex[®] cough*

NDC 82442-751-04

Children's Cough Relief

Dextromethorphan HBr 5 mg

Guaifenesin 100 mg

- Controls Cough
- Relieves Chest Congestion
- Breaks up Mucus

Ages 4 + years

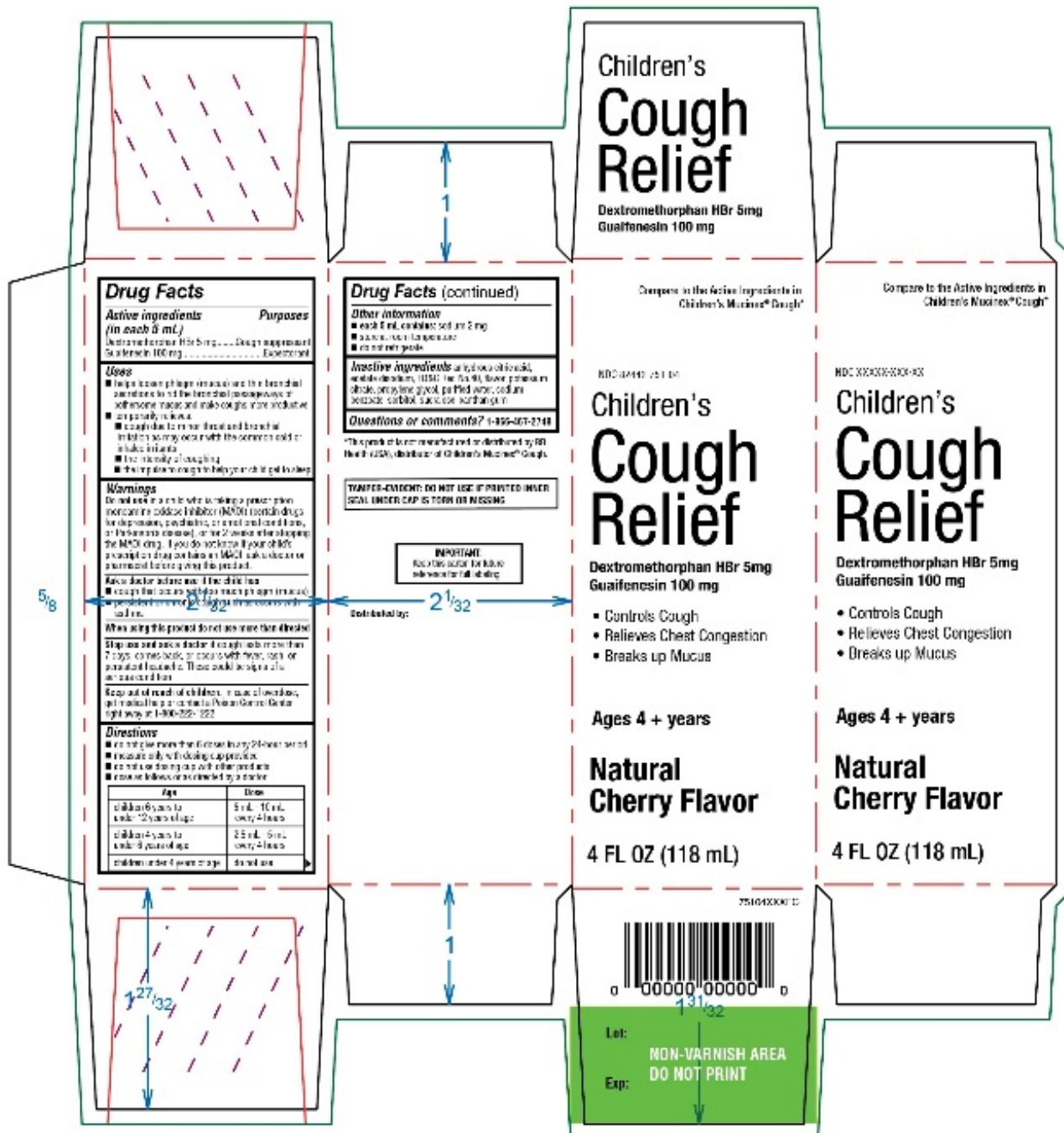
Natural Cherry Flavor

4 FL OZ(118 mL)

*This product is not manufactured or distributed by RB Health (USA), distributor of Children's Mucinex® Cough.

IMPORTANT: Keep this carton for future reference on full labeling.

Distributed by:



TARGET CHILDRENS COUGH RELIEF
dextromethorphan hydrobromide and guaifenesin liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82442-751
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	5 mg in 5 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	100 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
EDETATE SODIUM (UNII: MP1J8420LU)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
POTASSIUM CITRATE (UNII: EE90ONI6FF)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0K00R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	RED	Score	
Shape		Size	
Flavor	CHERRY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82442-751-04	1 in 1 CARTON	02/02/2024	
1		118 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 Drug	M012	02/02/2024	

