DRX CHOICE CHILDRENS STUFFY NOSE AND CHEST CONGESTION- guaifenesin and phenylephrine hcl solution
RARITAN PHARMACEUTICALS

DRx Choice Children's Stuffy Nose & Chest Congestion

Drug Facts

Active ingredients (in each 5 mL)

Guaifenesin, USP 100 mg

Phenylephrine HCl, USP 2.5 mg

Purposes

Expectorant
Nasal Decongestant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive
- temporarily relieves:
  - nasal congestion due to a cold
  - stuffy nose

Warnings

Do not usein 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

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- 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

- nervousness, dizziness or sleeplessness occur
- symptoms do not get better within 7 days or occur with fever
- cough lasts more than 7 days, comes back, or occurs with fever, rash, or persistent headache.

These could be signs of a serious illness.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away 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	10 mL every 4 hours
Children 4 years to under 6 years	5 mL every 4 hours
Children under 4 years	**do not use**

Other information

- **each 5 mL contains**: sodium 3 mg
- very low sodium
- store at room temperature
- do not refrigerate
- dosing cup provided

Inactive ingredients

anhydrous citric acid, edetate disodium, flavors, glycerin, propylene glycol, purified water, sodium benzoate, sodium citrate dihydrate, sorbitol solution, sucralose, xanthan gum.

Questions or Comments?

1-866-467-2748

Package Label-Principal Display Panel 4 FL OZ (118 mL Bottle)

*Compare to the active ingredients in Children’s Mucinex® Stuffy Nose & Chest Congestion

NDC 68163-748-04

DRx Choice®

Children's

Stuffy Nose & Chest Congestion

Guaifenesin (Expectorant)
Phenylephrine HCl (Nasal Decongestant)

Relieves: Stuffy Nose, Chest congestion, Breaks Up Mucus

Dosage Cup Included

Mixed Berry Flavor

Naturally & artificially flavored

For Ages 4 & Over

4 FL OZ (118 mL)

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

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

Manufactured by:

Raritan Pharmaceuticals

8 Joanna Court. East Brunswick

NJ 08816

DRX CHOICE CHILDRENS STUFFY NOSE AND CHEST CONGESTION

guaifenesin and phenylephrine hcl solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68163-748
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	100 mg in 5 mL

PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	2.5 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
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)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	pink	**Score**	
Shape		**Size**	
Flavor	BERRY	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68163-748-04	1 in 1 CARTON	04/24/2023	
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	04/24/2023	

Labeler - RARITAN PHARMACEUTICALS (127602287)

Revised: 9/2025 RARITAN PHARMACEUTICALS