DRX CHOICE CHILDRENS ALLERGY CHEWS- diphenhydramine hcl tablet, chewable
RARITAN PHARMACEUTICALS INC

DRx Choice children's allergy chews

Active ingredient (in each tablet)

Diphenhydramine HCl 12.5 mg

Purpose

Antihistamine

Uses

- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies.
    - runny nose
    - itchy, watery eyes
    - sneezing
    - itching of the nose or throat

Warnings

Do not use

- to make a child sleepy
- with any other product containing diphenhydramine, even one used on skin

Ask your doctor before use if you have

- glaucoma
- a breathing problem such as emphysema or chronic bronchitis
- trouble urinating due to an enlarged prostate gland

Ask a doctor or pharmacist before use if you are

taking sedatives or tranquilizers.

When using this product

- marked drowsiness may occur
- avoid alcoholic drinks
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery

- excitability may occur, especially in children

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

- chew one tablet completely at the onset of symptoms. Do not swallow tablets whole.
- Find right dose on chart below
- take every 4 to 6 hours, or as directed by a doctor
- do not take more than 6 doses in 24 hours

Age (Yr)	**Dose (chewable tablets)**
children under 2 years of age	do not use
children 2 to under 5 years of age	do not use unless directed by a doctor
children 6 to under 12 years of age	1 to 2 tablets
adults and children 12 years of age and over	2 to 4 tablets

Other information

- store below at room temperature. Avoid high humidity. Protect from light.

Inactive ingredients

citric acid, crospovidone, D&C Red No.30, dextrose, FD&C Blue No 1, flavors, magnesium stearate, maltodextrin, potassium citrate, silica, sodium polystyrene sulfonate, starch, sucralose

Questions or comments ?

1-866-467-2748

Principal Display Panel

NDC 68163-012-48

Compare to active ingredient in Children's Benadryl® Allergy Chewable Tablets

DRx Choice

children's allergy chews

diphenhydramine HCl, 12.5 mg/ antihistamine

For Relief of:

- Sneezing
- Runny nose
- Itchy throat or Nose
- Itchy, watery eyes

48 Chewable Tablets

Grape Flavor

4-6 HOURS/DOSE

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING

*This product is not manufactured or distributed by Johnson & Johnson Consumer INC, owner of the registered trademark Children's Benadryl® Allergy Chewable Tablets.

Manufactured by:

Raritan Pharmaceuticals

8 Joanna Court,

East Brunswick,

NJ 08816

IMPORTANT: Keep this carton for future reference on full labelling

DRX CHOICE CHILDRENS ALLERGY CHEWS

diphenhydramine hcl tablet, chewable

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68163-012
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	12.5 mg

Inactive Ingredients	
Ingredient Name	**Strength**
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
CROSPOVIDONE (120 .MU.M) (UNII: 68401960MK)	
D&C RED NO. 30 (UNII: 2S42T2808B)	
DEXTROSE, UNSPECIFIED FORM (UNII: IY9XDZ35W2)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
POTASSIUM CITRATE (UNII: EE90ONI6FF)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM POLYSTYRENE SULFONATE (UNII: 1699G8679Z)	
STARCH, CORN (UNII: O8232NY3SJ)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics

Color	PURPLE	**Score**	2 pieces
Shape	ROUND	**Size**	16mm
Flavor	GRAPE	**Imprint Code**	RP012
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68163-012-48	4 in 1 CARTON	07/04/2022	
1		12 in 1 BLISTER PACK; 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	07/04/2022	

Labeler - RARITAN PHARMACEUTICALS INC (127602287)

Revised: 11/2024 RARITAN PHARMACEUTICALS INC