

**ALLERGY RELIEF CHILDRENS- diphenhydramine hcl solution**  
**L.N.K. International, Inc.**

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**Quality Plus 44-015**

***Active ingredient (in each 5 mL)***

Diphenhydramine HCl 12.5 mg

***Purpose***

Antihistamine

***Uses***

- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:
  - runny nose
  - sneezing
  - itchy, watery eyes
  - 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 a doctor before use if the child has**

- glaucoma
- a breathing problem such as chronic bronchitis

**Ask a doctor or pharmacist before use if the child is**

taking sedatives or tranquilizers.

**When using this product**

- marked drowsiness may occur
- sedatives and tranquilizers may increase drowsiness
- excitability may occur, especially in children

**Keep out of reach of children.**

In case of overdose, get medical help or contact a Poison Control Center right away.

***Directions***

- do not take more than 6 doses in 24 hours
- mL = milliliter; FL OZ = fluid ounce
- use only enclosed dosing cup designed for use with this product. Do not use any other dosing device.
- find right dose on chart below
- take every 4 to 6 hours, or as directed by a doctor

<b>Age (yr)</b>	<b>Dose (mL)</b>
children under 2 years	do not use
children 2 to 5 years	do not use unless directed by a doctor
children 6 to 11 years	5 mL to 10 mL

***Other information***

- **each 5 mL contains:** sodium 5 mg
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- see end flap for expiration date and lot number
- protect from light

***Inactive ingredients***

anhydrous citric acid, D&C red #33, FD&C red #40, flavors, glycerin, high fructose corn syrup, purified water, sodium benzoate, sodium chloride, sodium citrate dihydrate, sucrose

***Questions or comments?***

**1-800-426-9391**

***Principal display panel***

**QUALITY  
+PLUS**

NDC 50844-015-19

Compare to active ingredient  
in Children's Benadryl® Allergy\*

**CHILDREN'S  
ALLERGY  
RELIEF**

**Diphenhydramine HCl 12.5 mg  
Antihistamine**

**Relieves**

- **Sneezing**
  - **Runny nose**
  - **Itchy, Watery eyes**
  - **Itchy throat**
- or nose**

**Cherry  
Flavor**

**8 FL OZ (237 mL)**

**ALCOHOL FREE**

**TAMPER EVIDENT: DO NOT USE IF IMPRINTED  
SAFETY SEAL UNDER CAP IS BROKEN OR MISSING**

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

50844                    ORG051901519

Distributed by  
**LNK INTERNATIONAL, INC.**

60 Arkay Drive  
Hauppauge, NY 11788  
USA



## ALLERGY RELIEF CHILDRENS

diphenhydramine hcl solution

### Product Information

**Product Type**

HUMAN OTC DRUG

**Item Code (Source)**

NDC:50844-015

<b>Route of Administration</b>	ORAL
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<b>Active Ingredient/Active Moiety</b>		
<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>DIPHENHYDRAMINE HYDROCHLORIDE</b> (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	12.5 mg in 5 mL

<b>Inactive Ingredients</b>	
<b>Ingredient Name</b>	<b>Strength</b>
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>D&amp;C RED NO. 33</b> (UNII: 9DBA0SBB0L)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>HIGH FRUCTOSE CORN SYRUP</b> (UNII: XY6UN3QB6S)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>TRISODIUM CITRATE DIHYDRATE</b> (UNII: B22547B95K)	
<b>SUCROSE</b> (UNII: C151H8M554)	

<b>Product Characteristics</b>			
<b>Color</b>	red	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	CHERRY	<b>Imprint Code</b>	
<b>Contains</b>			

<b>Packaging</b>				
<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:50844-015-19	1 in 1 CARTON	06/28/2024	
1		237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

<b>Marketing Information</b>			
<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC Monograph Drug	M012	06/28/2024	

**Labeler** - L.N.K. International, Inc. (038154464)

**Establishment**

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	manufacture(50844-015) , pack(50844-015)

Revised: 6/2024

L.N.K. International, Inc.