

CHILDRENS ALLERGY RELIEF- diphenhydramine hcl tablet, chewable
Topco Associates, LLC

TopCare 44-585

Active ingredient (in each chewable 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

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

Ask a doctor before use if you have

- glaucoma
- a breathing problem such as emphysema or chronic bronchitis
- difficulty in urination due to enlargement of the prostate gland

Ask a doctor or pharmacist before use if you are

taking sedatives or tranquilizers.

When using this product

- alcohol, sedatives, and tranquilizers may increase drowsiness
- use caution when driving a motor vehicle or operating machinery
- avoid alcoholic beverages
- marked drowsiness may occur
- 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.

Directions

- find right dose on chart below
- chew or crush tablets completely before swallowing; do not swallow tablets whole
- take every 4 to 6 hours, or as directed by a doctor
- do not take more than 6 times in 24 hours

| Age (yr) | Dose (chewable tablets) |
|---------------------------------------|--|
| 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 | 1 to 2 chewable tablets (12.5 mg to 25 mg) |
| adults and children 12 years and over | 2 to 4 chewable tablets (25 mg to 50 mg) |

Other information

- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN**
- store at 25°C (77°F); excursions permitted between 15°C-30°C (59°F-86°F)
- avoid high humidity
- see end flap for expiration date and lot number

Inactive ingredients

D&C red #27 aluminum lake, D&C red #30 aluminum lake, dextrans hydrated, ethylcellulose, FD&C blue #1 aluminum lake, flavor, hydroxypropyl cellulose, magnesium stearate, mannitol, stearic acid, sucralose, sucrose

Questions or comments?

1-888-423-0139

Principal Display Panel

+TopCare®
health

NDC 36800-585-44

COMPARE TO CHILDREN'S BENADRYL® CHEWABLES ACTIVE INGREDIENT*

children's
Allergy Relief

DIPHENHYDRAMINE HCl 12.5 mg
CHEWABLE TABLETS - ANTIHISTAMINE

**4-6
HOURS/
DOSE**

RELIEF OF:

- Sneezing
- Runny Nose
- Itchy, Watery Eyes
- Itchy Nose or Throat

**18 CHEWABLE
TABLETS**

Chew or crush tablets completely before swallowing.

For Ages 6 to 11 Years

**GRAPE
FLAVOR**

actual size

**TAMPER EVIDENT: DO NOT USE IF PACKAGE 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 Corporation,
owner of the registered trademark Children's Benadryl® Chewables.

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topcare@topco.com
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**QUALITY
GUARANTEED**

+TopCare®
health

**18 CHEWABLE
TABLETS**

children's
Allergy Relief

DIPHENHYDRAMINE HCl 12.5 mg
ANTIHISTAMINE

NDC 36800-585-44

+TopCare®
health

COMPARE TO CHILDREN'S
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ACTIVE INGREDIENT*

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CHEWABLE TABLETS - ANTIHISTAMINE



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**18 CHEWABLE
TABLETS**

Chew or crush tablets completely
before swallowing.
For Ages 6 to 11 Years

**GRAPE
FLAVOR** actual size

No Print / No Varnish
Lot no. & Exp. date



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

**✓ QUALITY
GUARANTEED**

Scan here for more
information or
call 1-888-423-0139



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50844 REV1218858544

Questions or comments? 1-888-423-0139

Inactive ingredients D&C red #27 aluminum lake, D&C
red #30 aluminum lake, dehydrated, ethylcellulose, FD&C
blue #1 aluminum lake, flavor, hydroxypropyl cellulose,
magnesium stearate, mannitol, stearic acid, sucrose, sucrose

Other information
■ TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS
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(59°-86°F) ■ avoid high humidity
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(12.5 mg to 25 mg)
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(25 mg to 50 mg)

Drug Facts (continued)

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

■ pregnant or breast-feeding, ask a health professional before use.
■ avoid alcohol or other sedatives, and tranquilizers may increase drowsiness
■ use caution when driving a motor vehicle or operating machinery
■ avoid alcoholic beverages ■ marked drowsiness may occur
■ excitability may occur, especially in children

■ Ask a doctor or pharmacist before use if you are taking sedatives
or tranquilizers.
■ difficulty in urination due to enlargement of the prostate gland
■ a breathing problem such as emphysema or chronic bronchitis
■ Ask a doctor before use if you have ■ glaucoma

■ with any other product containing diphenhydramine, even one
used on skin ■ to make a child sleepy
Do not use
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Warnings
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■ a breathing problem such as emphysema or chronic bronchitis
■ difficulty in urination due to enlargement of the prostate gland
■ use caution when driving a motor vehicle or operating machinery
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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

Active ingredient
Diphenhydramine HCl 12.5 mgAntihistamine
Purpose
Drug Facts
**KEEP OUTER PACKAGE FOR
COMPLETE PRODUCT INFORMATION**

CHILDRENS ALLERGY RELIEF

diphenhydramine hcl tablet, chewable

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:36800-585 |
| 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 |
|--|----------|
| D&C RED NO. 27 ALUMINUM LAKE (UNII: ZK64F7XSTX) | |
| D&C RED NO. 30 (UNII: 2S42T2808B) | |
| DEXTROSE MONOHYDRATE (UNII: LX22YL083G) | |
| ETHYLCELLULOSE, UNSPECIFIED (UNII: 7Z8S9VYZ4B) | |
| FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM) | |
| HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH) | |
| MAGNESIUM STEARATE (UNII: 70097M6I30) | |
| MANNITOL (UNII: 3OWL53L36A) | |
| STEARIC ACID (UNII: 4ELV7Z65AP) | |
| SUCRALOSE (UNII: 96K6UQ3ZD4) | |
| SUCROSE (UNII: C151H8M554) | |

Product Characteristics

| | | | |
|-----------------|--------|---------------------|----------|
| Color | purple | Score | no score |
| Shape | ROUND | Size | 12mm |
| Flavor | GRAPE | Imprint Code | 44;585 |
| Contains | | | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:36800-585-44 | 3 in 1 CARTON | 03/20/2009 | |
| 1 | | 6 in 1 BLISTER PACK; Type 0: Not a Combination Product | | |

Marketing Information

| Marketing | Application Number or Monograph | Marketing Start | Marketing End |
|-----------|---------------------------------|-----------------|---------------|
|-----------|---------------------------------|-----------------|---------------|

| Category | Citation | Date | Date |
|--------------------|----------|------------|------|
| OTC Monograph Drug | M012 | 03/20/2009 | |

Labeler - Topco Associates, LLC (006935977)

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|---------------------|
| LNK International, Inc. | | 038154464 | pack(36800-585) |

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|---------------------|
| LNK International, Inc. | | 832867837 | pack(36800-585) |

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|------------------------|
| LNK International, Inc. | | 832867894 | manufacture(36800-585) |

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|---------------------|
| LNK International, Inc. | | 868734088 | pack(36800-585) |

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|---------------------|
| LNK International, Inc. | | 967626305 | pack(36800-585) |