

**BENADRYL- diphenhydramine hydrochloride**  
**Johnson & Johnson Consumer Inc.**

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**Benadryl**

***Drug Facts***

**Active ingredient (in each tablet)**

Diphenhydramine HCl 25 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
- temporarily relieves these symptoms due to the common cold:
  - runny nose
  - sneezing

**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 you have**

- a breathing problem such as emphysema or chronic bronchitis
- glaucoma
- 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. (1-800-222-1222)

### Directions

- take every 4 to 6 hours, or as directed by a doctor
- do not take more than 6 times in 24 hours

adults and children 12 years and over	1 to 2 tablets
children 6 to under 12 years	1 tablet
children under 6 years	do not use

### Other information

- **each tablet contains:** calcium 15 mg
- store between 20-25°C (68-77°F). Protect from light.
- **do not use if carton is opened or carton tape and foil inner seal imprinted with "SAFETY SEAL®" are broken or missing**

### Inactive ingredients

carnauba wax, croscarmellose sodium, D&C red no. 27 aluminum lake, dibasic calcium phosphate, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate 80, titanium dioxide

### Questions or comments?

call **1-877-717-2824** (toll-free) or **215-273-8755** (collect)

### PRINCIPAL DISPLAY PANEL

**100+48 = 148**

**ULTRATABS®\***  
**TABLETS**

**Benadryl®**

**WORKS WHEN YOU  
NEED IT MOST™**

\*small tablet size

0 00000 00000 0  
FPO  
Exp:  
Lot:

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Abbott Laboratories  
Abbott Park, IL 60014  
U.S. Patent 5,412,111



100 CT.  
BOTTLE

**ON-THE-GO  
OR AT HOME**



48 CT.  
BLISTER

# RELIEF YOU CAN TRUST FOR ALL SEASONS

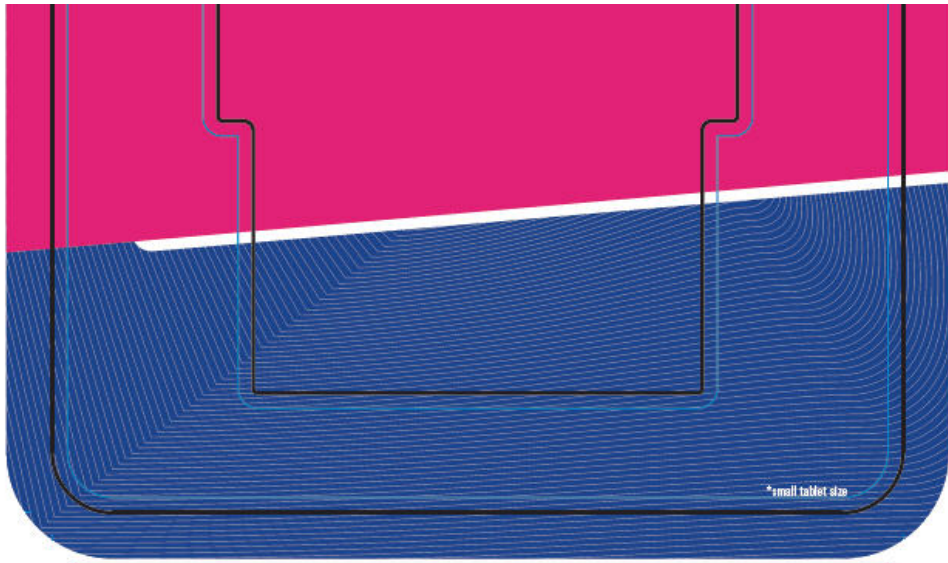
- 👉 Sneezing
- 👉 Runny Nose
- 👉 Itchy, Watery Eyes
- 👉 Itchy Throat



**100 + 48 = 148** ULTRATABS<sup>®</sup>  
TABLETS



**WORKS WHEN YOU  
NEED IT MOST™**



\*small tablet size

# Benadryl<sup>®</sup> ALLERGY

**EFFECTIVE ALLERGY RELIEF WHEN YOU NEED IT!<sup>®</sup>**

For allergy tips and information, visit [www.benadryl.com](http://www.benadryl.com)

## Allergy Tips



### Close the windows

Keep your windows closed to help keep allergens on the outside. If it gets warm, use an air-conditioner.



### Change your clothes

If you've worn it outside, don't wear it inside. That way you avoid spreading allergens around your home.



### Take off your shoes

Leave your shoes at the front door to avoid tracking dust and allergens into the home

NDC 50580-226-50

# Benadryl<sup>®</sup>

# ALLERGY

**Diphenhydramine HCl 25 mg**  
Antihistamine

**ULTRATABS<sup>®</sup>\***

\*small tablet size



actual  
size

**100 TABLETS**

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ibnal before usa

ison Control

door

2 tablets

tablet

not use

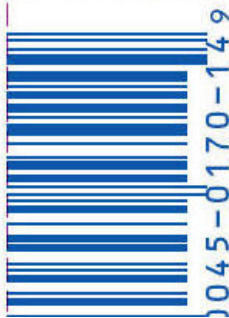
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taken or

mellrose sodium  
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NSUNER INC.

16 30057325



045-0170-149

**Important:** Read all product information before using. See this box for important information.

**Drug Facts**  
**Active ingredient (in each tablet)** Diphenhydramine HCl 25 mg.....Amlisilamine

**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  
 temporarily relieves these symptoms due to the common cold:  
 ■ runny nose ■ sneezing

**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 you have**  
 ■ a breathing problem such as emphysema or chronic bronchitis  
 ■ glaucoma ■ 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

**Drug Facts (continued)**  
 ■ be careful when driving a motor vehicle or operating machinery if drowsiness occurs, especially in children  
 ■ if pregnant or breast-feeding, ask a health professional for advice before using. Keep out of reach of children.  
 In case of overdose, get medical help or contact a Poison Control Center right away. (1-800-232-7272)

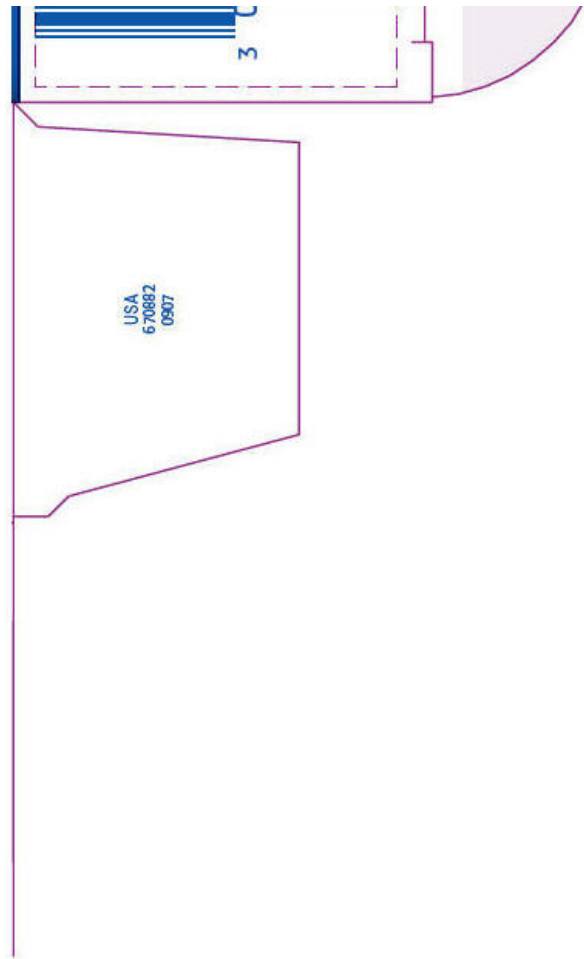
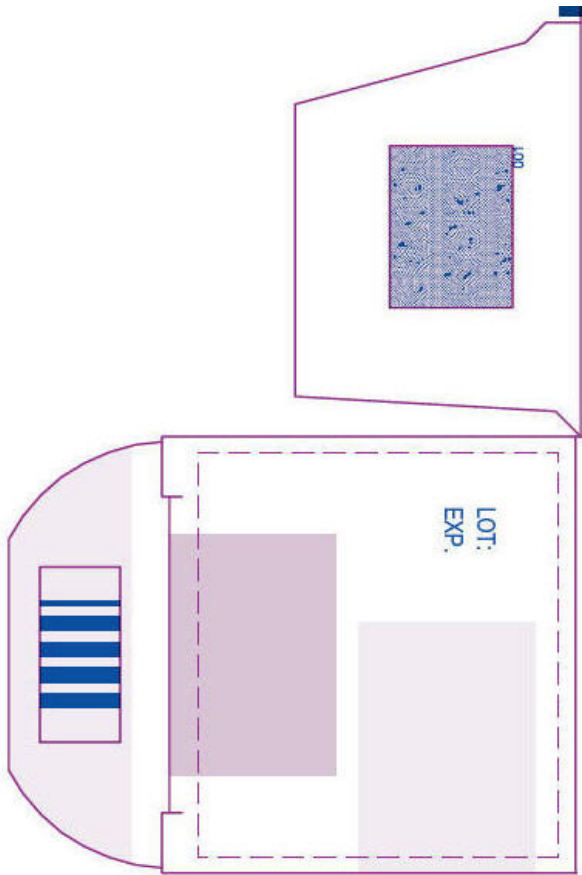
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 adults and children 12 years and over 1 to 2 tablets  
 children 6 to under 12 years 1 to 2 tablets  
 children under 6 years do not use

**Other information**  
 ■ each tablet contains: calcium 15 mg  
 ■ store between 20°-25°C (68°-77°F). Protect from light  
 ■ do not use if cap is opened or cap seal is missing  
 ■ seal imprinted with "SAFETY SEAL" are missing

**Inactive ingredients:** carmelumina, croscellan D&G, red no. 27 aluminum lake, dibasic calcium phosphate, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate 80, titanium dioxide

**Questions or comments?** call 1-877-717-7177 or 215-273-8735 (toll-free)

Active ingredient made in Japan  
 Distributed by: **JOHNSON & JOHNSON**  
 McKel Consumer Healthcare Division  
 Fort Washington, PA 19084 USA ©J&JCI 20



## BENADRYL

diphenhydramine hydrochloride kit

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:50580-370
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### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50580-370-01	1 in 1 PACKAGE	06/12/2017	

### Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE	100
Part 2	4 BLISTER PACK	48

### Part 1 of 2

## BENADRYL

diphenhydramine hydrochloride tablet, film coated

## Product Information

**Route of Administration** ORAL

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>DIPHENHYDRAMINE HYDROCHLORIDE</b> (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg

## Inactive Ingredients

Ingredient Name	Strength
<b>CARNAUBA WAX</b> (UNII: R12CBM0EIZ)	
<b>CROSCARMELLOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>D&amp;C RED NO. 27 ALUMINUM LAKE</b> (UNII: ZK64F7XSTX)	
<b>DIBASIC CALCIUM PHOSPHATE DIHYDRATE</b> (UNII: O7TSZ97GEP)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>POLYSORBATE 80</b> (UNII: 6OZP39ZG8H)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	

## Product Characteristics

<b>Color</b>	pink	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	11mm
<b>Flavor</b>		<b>Imprint Code</b>	B;WL;25
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		1 in 1 CARTON		
1		100 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	06/04/2012	

## Part 2 of 2

### BENADRYL

diphenhydramine hydrochloride tablet, film coated

#### Product Information

**Route of Administration** ORAL

#### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII: 8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg

#### Inactive Ingredients

Ingredient Name	Strength
CARNAUBA WAX (UNII: R12CBM0EIZ)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
D&C RED NO. 27 ALUMINUM LAKE (UNII: ZK64F7XSTX)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

#### Product Characteristics

<b>Color</b>	pink	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	11mm
<b>Flavor</b>		<b>Imprint Code</b>	B;WL;25
<b>Contains</b>			

#### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		4 in 1 CARTON		
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
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OTC Monograph Drug	M012	06/04/2012	
<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	06/12/2017	

**Labeler** - Johnson & Johnson Consumer Inc. (878046358)

Revised: 1/2024

Johnson & Johnson Consumer Inc.