

**BENADRYL- diphenhydramine hydrochloride tablet, film coated**  
**Mechanical Servants, LLC**

*Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.*

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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 pouch is torn or damaged**

### 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)

Distributed by:

**JOHNSON & JOHNSON CONSUMER INC.**

McNeil Consumer Healthcare Division

Fort Washington, PA 19034 USA

Repackaged and distributed by Convenience Valet<sup>®</sup>, Melrose Park, IL 60160

### PRINCIPAL DISPLAY PANEL - 25 mg Tablet Pouch Package

**Children's**

**Miracle Network**

Hospitals<sup>®</sup>

minimum annual

**\$250,000**

**Donation**

See details on back.

**Benadryl®*****ALLERGY*****Diphenhydramine HCl 25 mg | Antihistamine**

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

**ULTRATABS®\***

\*small tablet size

**2 Pouches of 2 Tablets each**

www.childrensmiraclenetworkhospitals.org  
Convenience Valet donates \$.01  
to Children's Miracle  
Network Hospitals for the  
purchase of this package.

Minimum Donation \$250,000 (per year)  
cvalet.com

Do not use if  
package or pouch  
is opened

**Benadryl**<sup>®</sup>  
**ALLERGY**

Diphenhydramine HCl 25 mg | Antihistamine



Important: Read all product information before using.  
Keep this package for important information.

### Drug Facts

Active ingredient (in each tablet)	Purpose
Diphenhydramine HCl 25 mg	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

The Children's Miracle Network Hospitals name and logo are used with permission which in no way constitutes endorsement, express or implied, of any commercial product or company.

OPEN PANEL FOR CONTINUED DRUG FACTS

Repackaged and distributed by Convenience Valet<sup>®</sup>, Melrose Park, IL 60160

1007.04.17.FB / CBB-MP-DFB-1007

minimum annual  
**\$250,000**  
Donation  
See details on back.



Children's  
Miracle Network  
Hospitals<sup>®</sup>

**Benadryl**<sup>®</sup>

**ALLERGY**

Diphenhydramine HCl 25 mg | Antihistamine

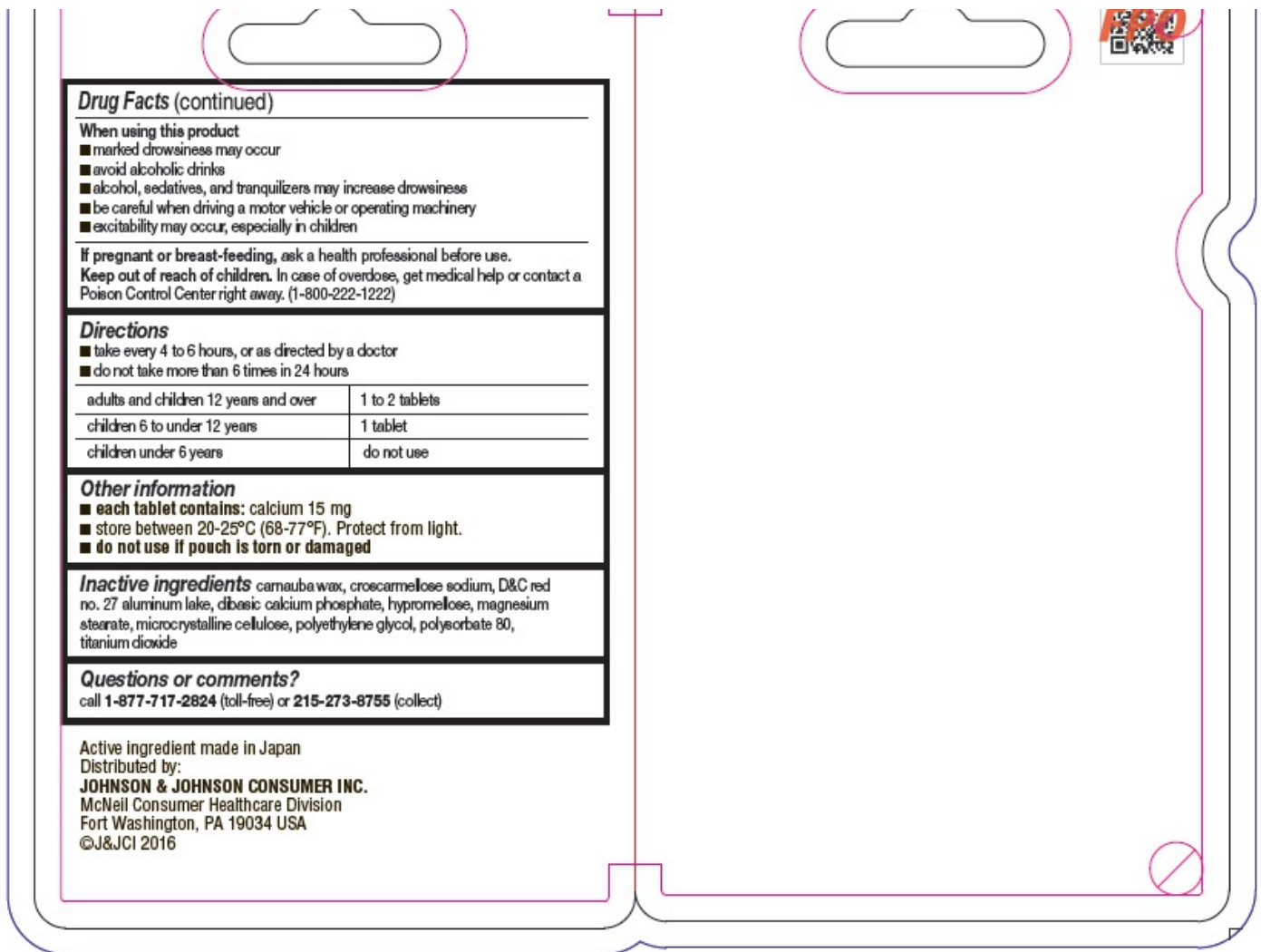
- ✓ Sneezing
- ✓ Itchy, Watery Eyes
- ✓ Runny Nose
- ✓ Itchy Throat

ULTRATABS<sup>®</sup>\*



\*small tablet size

2 Pouches of 2 Tablets each



## PRINCIPAL DISPLAY PANEL-25mg Tablet Pouch Package 6ct

**Benadryl** ®

ALLERGY

**Diphenhydramine HCl 25 mg** | Antihistamine

Sneezing

Itchy, Watery Eyes

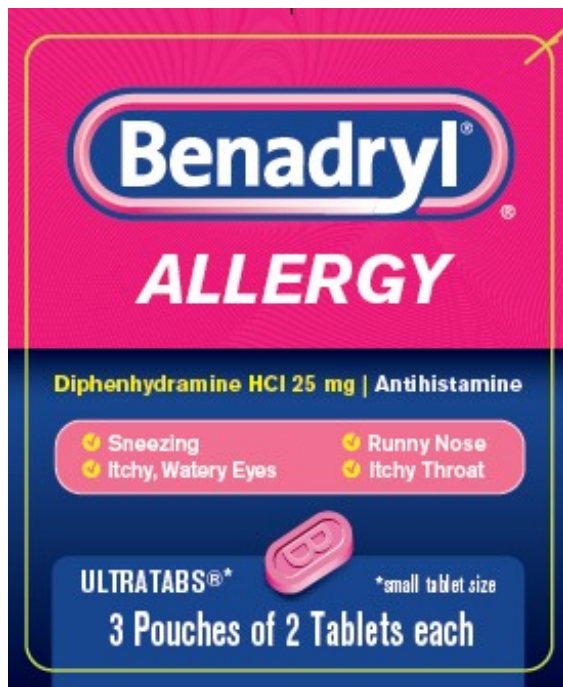
Runny Nose

Itchy Throat

ULTRATABS ®\*

\*small tablet size

**3 POUCHES OF 2 TABLETS EACH**



**PRINCIPAL DISPLAY PANEL-25mg Tablet Pouch 25 Box**

**Benadryl ®**

ALLERGY

**Diphenhydramine HCl 25 mg | Antihistamine**

Sneezing  
Itchy, Watery Eyes  
Runny Nose  
Itchy Throat

ULTRATABS ®\*  
\*small tablet size

**25 POUCHES OF 2 TABLETS EACH**



**PRINCIPAL DISPLAY PANEL-25mg Tablet Pouch 30 Box**

**Benadryl ®**

ALLERGY

**Diphenhydramine HCl 25 mg | Antihistamine**

Sneezing  
Itchy, Watery Eyes  
Runny Nose  
Itchy Throat

ULTRATABS ®\*  
\*small tablet size

**30 POUCHES OF 2 TABLETS EACH**





## BENADRYL

diphenhydramine hydrochloride tablet, film coated

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:29485-6931
<b>Route of Administration</b>	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>CROSCARMELOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>D&amp;C RED NO. 27</b> (UNII: 2LRS185U6K)	



<b>ALUMINUM OXIDE</b> (UNII: LMI26O6933)	
<b>ANHYDROUS DIBASIC CALCIUM PHOSPHATE</b> (UNII: L11K75P92J)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I3O)	
<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	NDC:29485-6931-6	3 in 1 BLISTER PACK	02/06/2017	
1		2 in 1 POUCH; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	02/06/2017	

## BENADRYL

diphenhydramine hydrochloride tablet, film coated

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:29485-7963
<b>Route of Administration</b>	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
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<b>CARNAUBA WAX</b> (UNII: R12CBM0EIZ)
<b>CROSCARMELOSE SODIUM</b> (UNII: M28OL1HH48)
<b>D&amp;C RED NO. 27</b> (UNII: 2LRS185U6K)
<b>ALUMINUM OXIDE</b> (UNII: LMI26O6933)
<b>ANHYDROUS DIBASIC CALCIUM PHOSPHATE</b> (UNII: L11K75P92J)
<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	NDC:29485-7963-2	25 in 1 BOX	07/10/2020	
1		2 in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:29485-7963-3	30 in 1 BOX	10/19/2015	
2		2 in 1 POUCH; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	10/19/2015	

## BENADRYL

diphenhydramine hydrochloride tablet, film coated

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:29485-1007
<b>Route of Administration</b>	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>CROSCARMELOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>D&amp;C RED NO. 27</b> (UNII: 2LRS185U6K)	
<b>ALUMINUM OXIDE</b> (UNII: LMI26O6933)	
<b>ANHYDROUS DIBASIC CALCIUM PHOSPHATE</b> (UNII: L11K75P92J)	
<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	NDC:29485-1007-4	2 in 1 PACKAGE	09/01/2008	
1		2 in 1 POUCH; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	09/01/2008	

**Labeler** - Mechanical Servants, LLC (005530951)

### Establishment

Name	Address	ID/FEI	Business Operations
Mechanical Servants, LLC		005530951	relabel(29485-1007, 29485-6931, 29485-7963) , repack(29485-1007, 29485-6931, 29485-7963)

## Establishment

Name	Address	ID/FEI	Business Operations
Johnson & Johnson Consumer Inc., McNeil Consumer Healthcare Divison		878046358	manufacture(29485-1007, 29485-6931, 29485-7963) , analysis(29485-1007, 29485-6931, 29485-7963) , label(29485- 1007, 29485-6931, 29485-7963) , pack(29485-1007, 29485-6931, 29485-7963)

Revised: 12/2021

Mechanical Servants, LLC