BENADRYL- diphenhydramine hydrochloride tablet, film coated
Johnson & Johnson Consumer Inc., McNeil Consumer Healthcare Division

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.

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 if blister unit is broken**

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

NDC 50580-226-51

Benadryl ®

ALLERGY

Diphenhydramine HCl 25mg | Antihistamine

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

ULTRATABS ®*

*small tablet size

actual size

24 TABLETS

NDC 50580-226-51

Benadryl®

ALLERGY

Diphenhydramine HCl 25mg | Antihistamine

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

EFFECTIVE ALLERGY RELIEF WHEN YOU NEED IT!®

Benadryl® ALLERGY

30037905

CHILD-RESISTANT PACKAGING

Benadryl® ALLERGY

Active ingredient made in Japan
Distributed by:
JOHNSON & JOHNSON CONSUMER INC.
McNeil Consumer Healthcare Division
Fort Washington, PA 19034 USA
©J&JCI 2016

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

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Benadryl® ALLERGY

30037905

BENADRYL

diphenhydramine hydrochloride tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50580-226
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: 3WQ0SDW1A)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	pink	**Score**	no score
Shape	OVAL	**Size**	11mm
Flavor		**Imprint Code**	B;WL;25
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50580-226-50	1 in 1 CARTON	06/04/2012	
1		100 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:50580-226-51	2 in 1 CARTON	06/04/2012	
2		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:50580-226-53	2 in 1 POUCH; Type 0: Not a Combination Product	06/04/2012	
4	NDC:50580-226-54	60 in 1 CARTON	07/27/2015	
4		2 in 1 POUCH; Type 0: Not a Combination Product		
5	NDC:50580-226-62	4 in 1 CARTON	01/02/2017	
5		2 in 1 POUCH; Type 0: Not a Combination Product		
6	NDC:50580-226-52	4 in 1 CARTON	06/04/2012	
6		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
7	NDC:50580-226-56	3 in 1 PACKAGE	02/01/2013	
7		4 in 1 CARTON		
7		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 final	part341	09/01/2008	

Labeler - Johnson & Johnson Consumer Inc., McNeil Consumer Healthcare Division (878046358)

Revised: 10/2021 Johnson & Johnson Consumer Inc., McNeil Consumer Healthcare Division