BENADRYL ALLERGY EXTRA STRENGTH- diphenhydramine hydrochloride tablet, film coated
Johnson & Johnson Consumer Inc.

Benadryl ®

Drug Facts

Active ingredient (in each tablet)

Diphenhydramine HCl 50 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 tablet
children under 12 years	do not use

Other information

- each tablet contains: **calcium 35 mg**
- store between 20-25°C (68-77°F). Protect from light.
- **do not use if blister unit is torn or broken**

Inactive ingredients

carnauba wax, croscarmellose sodium, 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

NEW NDC 50580-533-24

Benadryl ®

ALLERGY

EXTRA STRENGTH

Diphenhydramine HCl 50mg | Antihistamine

✓ **Sneezing** ✓ **Runny Nose**

✓ **Itchy, Watery Eyes** ✓ **Itchy Throat**

actual size

24 TABLETS

NEW

NDC 50580-533-24

Benadryl®

ALLERGY

EXTRA STRENGTH

Diphenhydramine HCl 50mg | Antihistamine

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

EFFECTIVE ALLERGY RELIEF WHEN YOU NEED IT!®

Benadryl® ALLERGY EXTRA STRENGTH

300505549

Active ingredient made in Japan
Distributed by:
JOHNSON & JOHNSON CONSUMER INC.
McNeil Consumer Healthcare Division
Fort Washington, PA 19034 USA
©J&JCI 2021 www.benadryl.com

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

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Benadryl® ALLERGY EXTRA STRENGTH

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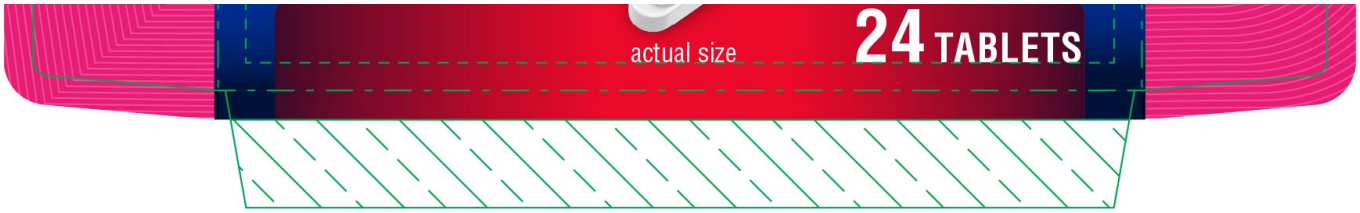

BENADRYL ALLERGY EXTRA STRENGTH

diphenhydramine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50580-533
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	50 mg

Inactive Ingredients

Ingredient Name	Strength
CARNAUBA WAX (UNII: R12CBM0EIZ)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
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	white	**Score**	no score
Shape	OVAL	**Size**	14mm
Flavor		**Imprint Code**	B;50MG
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50580-533-24	2 in 1 CARTON	07/05/2022	
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
OTC Monograph Drug	M012	07/05/2022	

Labeler - Johnson & Johnson Consumer Inc. (878046358)

Revised: 1/2024 Johnson & Johnson Consumer Inc.