

BANOPHEN- diphenhydramine hcl tablet, film coated
Proficient Rx LP

Rite Aid 44-329

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:
 - o runny nose
 - o itchy, watery eyes
 - o sneezing
 - o itching of the nose or throat
- temporarily relieves these symptoms due to the common cold:
 - o runny nose
 - o sneezing

Warnings

Do not use

- 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
- 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

- marked drowsiness may occur
- avoid alcoholic beverages

- alcohol, sedatives, and tranquilizers may increase drowsiness
- use caution 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.

Directions

- **do not take more than directed**
- 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 30 mg
- **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)
- protect from moisture
- see end flap for expiration date and lot number

Inactive ingredients

corn starch, D&C red #27 aluminum lake, dibasic calcium phosphate dihydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, silicon dioxide, stearic acid, talc, titanium dioxide

Questions or comments?

(800)-616-2471

Principal Display Panel

NDC 71205-611-20

Compare to the active ingredient in Benadryl® Allergy ULTRATAB® Tablets*

Banophen
Diphenhydramine HCl
25 mg
Antihistamine / Allergy Relief

Relieves
Sneezing, Runny Nose,
Itchy Throat and
Itchy, Watery Eyes

Actual Size

20 Minitabs

*This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark Benadryl® Allergy ULTRATAB® Tablets.
50844 REV1220M32908

Rev. 03/21 M-17 Re-order No. 250050

Distributed by:

MAJOR® PHARMACEUTICALS

17177 N Laurel Park Drive, Suite 233 Livonia, MI 48152 USA

Repackaged by:

PROFICIENT RX LP

Thousand Oaks, CA 91320

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

ProficientRx

Scan Here

NDC 71205-611-20

Packaged By: Proficient Rx LP
Thousand Oaks, CA 91320

Diphenhydramine HCl 25mg
#20 Tablets SN# MASTER
Lot #:00000 Exp:00/00/00
NDC 71205-611-20

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NDC 71205-611-20

GTIN: 00371205611200
SN# MASTER
Exp. 00/00/00
Lot #:00000

3
71205 61120 0

Diphenhydramine HCl 25mg
#20 Tablets

Each tablet contains: Diphenhydramine HCl 25 mg
Antihistamine

Pink, oval shaped, unscored tablet with imprint code "44 329"

Product ID: QD061120

Dist. By: MAJOR® PHARMACEUTICALS 17177 N Laurel Park Drive, Suite 233 Livonia, MI 48152 USA

Store at 25°C (77°F)

Keep medication out of the reach of children

BANOPHEN

diphenhydramine hcl tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71205-611(NDC:0904-5551)
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
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
D&C RED NO. 27 ALUMINUM LAKE (UNII: ZK64F7XSTX)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
STARCH, CORN (UNII: O8232NY3SJ)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
TALC (UNII: 7SEV7J4R1U)	

Product Characteristics

Color	pink	Score	no score
Shape	OVAL	Size	11mm
Flavor		Imprint Code	44;329
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71205-611-20	20 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/11/2021	
2	NDC:71205-611-30	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/11/2021	
3	NDC:71205-611-60	60 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/11/2021	
4	NDC:71205-611-90	90 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/11/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	03/02/1990	

Labeler - Proficient Rx LP (079196022)

Establishment

Name	Address	ID/FEI	Business Operations
Proficient Rx LP		079196022	REPACK(71205-611) , RELABEL(71205-611)

Revised: 12/2023

Proficient Rx LP