COMPLETE ALLERGY- diphenhydramine hcl tablet, film coated Chain Drug Consortium

Premier Value 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:
 - runny nose
 - itchy, watery eyes
 - sneezing
 - 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
- difficulty in urination due to enlargement of the prostate gland
- glaucoma

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

Other information

- each tablet contains: calcium 30 mg
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-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?

1-800-426-9391

Principal Display Panel

Premier Value®

*COMPARE TO THE ACTIVE INGREDIENT IN BENADRYL® ALLERGY ULTRATAB® TABLETS

Complete Allergy

Diphenhydramine HCl, 25 mg

ANTIHISTAMINE

Allergy relief for:

SneezingRunny nose

• Itchy, watery eyes • Itchy throat

actual size

24 Tablets

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

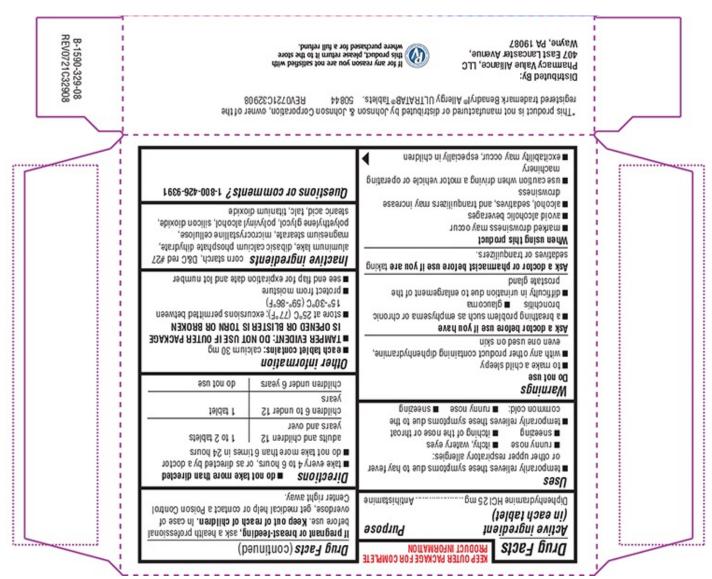
*This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the

registered trademark Benadryl® Allergy ULTRATAB® Tablets. 50844 REV0721C32908

Distributed By: Pharmacy Value Alliance, LLC 407 East Lancaster Avenue, Wayne, PA 19087

If for any reason you are not satisfied with this product, please return it to the store where purchased for a full refund.





Premier Value 44-329

COMPLETE ALLERGY

diphenhydramine hcl tablet, film coated

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68016-641	
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
STARCH, CORN (UNII: O8232NY3SJ)	

D&C RED NO. 27 ALUMINUM LAKE (UNII: ZK64F7XSTX)

DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)

MAGNESIUM STEARATE (UNII: 70097M6I30)

MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)

POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)

POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)

SILICON DIOXIDE (UNII: ETJ7Z6XBU4)

STEARIC ACID (UNII: 4ELV7Z65AP)

TALC (UNII: 7SEV7J4R1U)

TITANIUM DIOXIDE (UNII: 15FIX9V2JP)

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

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016- 641-24	2 in 1 CARTON	03/02/1990	
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:68016- 641-10	1 in 1 CARTON	03/02/1990	
2		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:68016- 641-00	1 in 1 CARTON	03/02/1990	
3		200 in 1 BOTTLE, PLASTIC; 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	03/02/1990		

Labeler - Chain Drug Consortium (101668460)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(68016-641)

Establishment			
Name	Address	ID/FEI	Business Operations

	manufacture	(68016-641).	pack(68016-641)	
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Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(68016-641)

832867837

LNK International, Inc.

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(68016-641)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(68016-641)

Revised: 10/2023 Chain Drug Consortium