WAL-DRYL ALLERGY- diphenhydramine hcl tablet, film coated Walgreen Company

Walgreens 44-329-Wal-Dryl-Delisted

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
- 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 (1-800-222-1222) right away.

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

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°-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, dicalcium phosphate, 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

Walgreens

NDC 0363-0329-08

Compare to Renadryl®

Allergy Ultratab[®] Tablets active ingredient^{††}

Wal-Dryl[®] ALLERGY

DIPHENHYDRAMINE HCl 25 mg / ANTIHISTAMINE

MINI TABS

- Relief of runny nose, sneezing, itchy throat & itchy, watery eyes
- Easy-to-swallow mini tabs

24

COATED MINI TABS

Actual Size

NOT FOR HOUSEHOLDS WITH YOUNG CHILDREN

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

DOES NOT CONTAIN GLUTEN

[†]Walgreens Pharmacist Survey *Walgreens* PHARMACIST RECOMMENDED[†]

Trusted since 1901[™]

Health expertise you rely on._{TM}

⁺⁺This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark Benadryl[®] Allergy Ultratab[®] Tablets.

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50844 REV1016A32908







diphenhydramine hcl tablet	, film coated					
Product Information						
Product Type	HUMAN OTC	DRUG	Item Code (S	Source)	NDC:036	3-0329
Route of Administration	ORAL	2.100		Jource,		
Route of Administration	OTAL					
Active Ingredient/Activ	e Moiety					
•	redient Nam	e		Basis of Str	renath	Strengt
DIPHENHYDRAMINE HYDROCH (DIPHENHYDRAMINE - UNII:8GTS8	ILORIDE (UNII:			DIPHENHYDRAMIN HYDROCHLORIDE	-	25 mg
Inactive Ingredients						
	Ingred	ient Name			S	trength
STARCH, CORN (UNII: 08232NY	3SJ)					
D&C RED NO. 27 ALUMINUM I	LAKE (UNII: ZK6	4F7XSTX)				
ANHYDROUS DIBASIC CALCIU	М РНОЅРНАТЕ	(UNII: L11K75	P92J)			
MAGNESIUM STEARATE (UNII:	70097M6I30)					
MICROCRYSTALLINE CELLULO	SE (UNII: OP1R	32D61U)				
POLYETHYLENE GLYCOL, UNS	SPECIFIED (UNI	I: 3WJQ0SDW1A	4)			
POLYVINYL ALCOHOL, UNSPE	CIFIED (UNII: 53	32B59J990)				
SILICON DIOXIDE (UNII: ETJ7Z6	SXBU4)					
STEARIC ACID (UNII: 4ELV7Z65/	AP)					
TALC (UNII: 7SEV7J4R1U)						
TITANIUM DIOXIDE (UNII: 15FIX	(9V2JP)					
Product Characteristic	s					
Color	pink	ink Score no score				
Shape	OVAL Size 11mm			.mm	n	
Flavor	Imprint Code 44;329					
Contains						
Packaging						
	Package De	scription	Ma	rketing Start Date		ting End Date

Ŧ	0329-08		02/02/1990	00/00/2025
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:0363- 0329-12	1 in 1 CARTON	03/02/1990	08/08/2025
2		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:0363- 0329-22	4 in 1 CARTON	03/02/1990	08/08/2025
3		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
4	NDC:0363- 0329-51	365 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/02/1990	08/08/2025
5	NDC:0363- 0329-78	600 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/02/1990	08/08/2025
Μ	arketing	Information		
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
	OTC Monograph Drug M012		03/02/1990	08/08/2025

Labeler - Walgreen Company (008965063)

Establishment					
Name	Address	ID/FEI	Business Operations		
LNK International, Inc.		038154464	pack(0363-0329)		

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(0363-0329) , pack(0363-0329)

Establishment					
Name	Address	ID/FEI	Business Operations		
LNK International, Inc.		832867894	manufacture(0363-0329)		
Establishment					

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(0363-0329)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(0363-0329)

Revised: 10/2023

Walgreen Company