WAL-SLEEP Z- diphenhydramine hcl tablet, film coated
Walgreen Company

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.

Walgreens 44-672 Delisted

Active ingredient (in each caplet)

Diphenhydramine HCl 25 mg

Purpose

Nighttime sleep-aid

Uses

- for relief of occasional sleeplessness
- reduces time to fall asleep if you have difficulty falling asleep

Warnings

Do not use

- for children under 12 years of age
- 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

avoid alcoholic beverages.

Stop use and ask a doctor if

sleeplessness persists continuously for more than 2 weeks. Insomnia may be a symptom of a serious underlying medical illness.

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

- **do not take more than directed**
- adults and children 12 years and over: take 2 caplets (50 mg) at bedtime if needed, or as directed by a doctor
- children under 12 years: do not use

Other information

- **each caplet contains:** calcium 65 mg
- avoid excessive heat (greater than 100°F) or humidity
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- see end flap for expiration date and lot number

Inactive ingredients

corn starch, D&C red #27 aluminum lake, dibasic calcium phosphate dihydrate, FD&C blue #2 aluminum lake, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, silicon dioxide, talc, titanium dioxide

Questions or comments?

1-800-426-9391

Principal display panel

Walgreens

Compare to VICKS®
ZzzQuil® active ingredient††

NDC 0363-0672-51

Wal-Sleep Z®

DIPHENHYDRAMINE HCl 25 mg /
NIGHTTIME SLEEP AID

NIGHTTIME CAPLETS

NOT FOR TREATING
PAIN, COLD OR FLU

365 CAPLETS

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

Walgreens Pharmacist Recommended
Walgreens Pharmacist Survey
††This product is not manufactured or distributed by Procter & Gamble, owner of the registered trademark Vicks® ZzzQuil®.

50844 REV0419A67251 REV0619

DISTRIBUTED BY:
WALGREEN CO.
200 WILMOT RD.
DEERFIELD, IL 60015
100% SATISFACTION GUARANTEED
walgreens.com

Walgreens 44-672

WAL-SLEEP Z
diphenhydramine hcl tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0363-0672
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)	
FD&C BLUE NO. 2--ALUMINUM LAKE (UNII: 4AQJ3LG584)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	purple	**Score**	no score
Shape	OVAL	**Size**	14mm
Flavor		**Imprint Code**	44;672
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-0672-51	365 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/24/2015	
2	NDC:0363-0672-21	1 in 1 CARTON	07/24/2015	10/14/2024
2		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:0363-0672-19	1 in 1 CARTON	07/24/2015	07/15/2019
3		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		
4	NDC:0363-0672-12	1 in 1 PACKAGE	07/24/2015	11/27/2020
4		100 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 final	part338	07/24/2015	10/14/2024

Labeler - Walgreen Company (008965063)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(0363-0672)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	pack(0363-0672)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(0363-0672) , pack(0363-0672)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(0363-0672)

Revised: 11/2022 Walgreen Company