DIPHENHYDRAMINE HCL MAXIMUM STRENGTH- diphenhydramine hcl tablet Bi-Mart

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Drug Facts

Active ingredient (in each caplet)

Diphenhydramine HCl 50 mg

Purpose

Nighttime sleep-aid

Use

helps reduce difficulty falling asleep

Warnings

Do not use

- in children under 12 years of age
- with any other product containing diphenhydramine, even one used on skin
- with other antihistamines

Ask a doctor before use if you have

- glaucoma
- a breathing problem such as emphysema or chronic bronchitis
- 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

- avoid alcoholic beverages
- be careful when driveing a motor vehicle or operating machinery

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 accidental overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

- **adults and children 12 years of age and over:** take 1 caplet at bedtime if needed, or as directed by your doctor
- children under 12 years: do not use

Other information

- **each caplet contains:**calcium 50 mg
- store at room temperature
- protect from moisture

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, dicalcium calcium phosphate dihydrate, FD&C blue #1 aluminum lake, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, silicon dioxide, stearic acid, talc, titanium dioxide, yellow iron oxide

Questions?

call toll free **1-844-912-4012**

Distributed by:

BI-MART

Eugene, OR 97402

Principal Display Panel

DIPHENHYDRAMINE HCL MAXIMUM STRENGTH

diphenhydramine hcl tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37835-852
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
FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
ANHYDROUS DIBASIC CALCIUM PHOSPHATE (UNII: L11K75P92J)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYVINYL ALCOHOL (UNII: 532B59J990)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	

Product Characteristics

Color	blue	**Score**	no score
Shape	OVAL (caplet)	**Size**	15mm
Flavor		**Imprint Code**	44;687
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37835-852-50	50 in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC Monograph Drug	M010	05/01/2025	

Labeler - Bi-Mart (027630078)

Registrant - Bi-Mart (027630078)

Establishment

Name	Address	ID/FEI	Business Operations
L.N.K. International, Inc.		038154464	manufacture(37835-852)

Revised: 5/2025 Bi-Mart