

**DIPHENHYDRAMINE HYDROCHLORIDE 50MG- diphenhydramine hydrochloride
50mg capsule, liquid filled
AMZ789 LLC**

diphenhydramine HCl 50mg softgels

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Use

For relief of occasional sleeplessness

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Warnings

for children under 12 years of age

with any other products containing diphenhydramine, even one used on skin

Ask a doctor or pharmacist before use if

you have a breathing problem such as emphysema or chronic bronchitis, glaucoma, trouble urinating due to an enlarged prostate gland

you are taking sedatives or tranquilizers

When using this product

avoid alcoholic drinks

Stop use and ask a doctor if

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

If pregnant or breast-feeding

ask a health professional before use

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

Direction

adults and children 12 years and over: 1 softgel at bedtime if needed, or as directed by a doctor

FD&C blue #1, gelatin, glycerin, polyethylene glycol, purified water, sorbitol sorbitan solution, white ink

RxZell

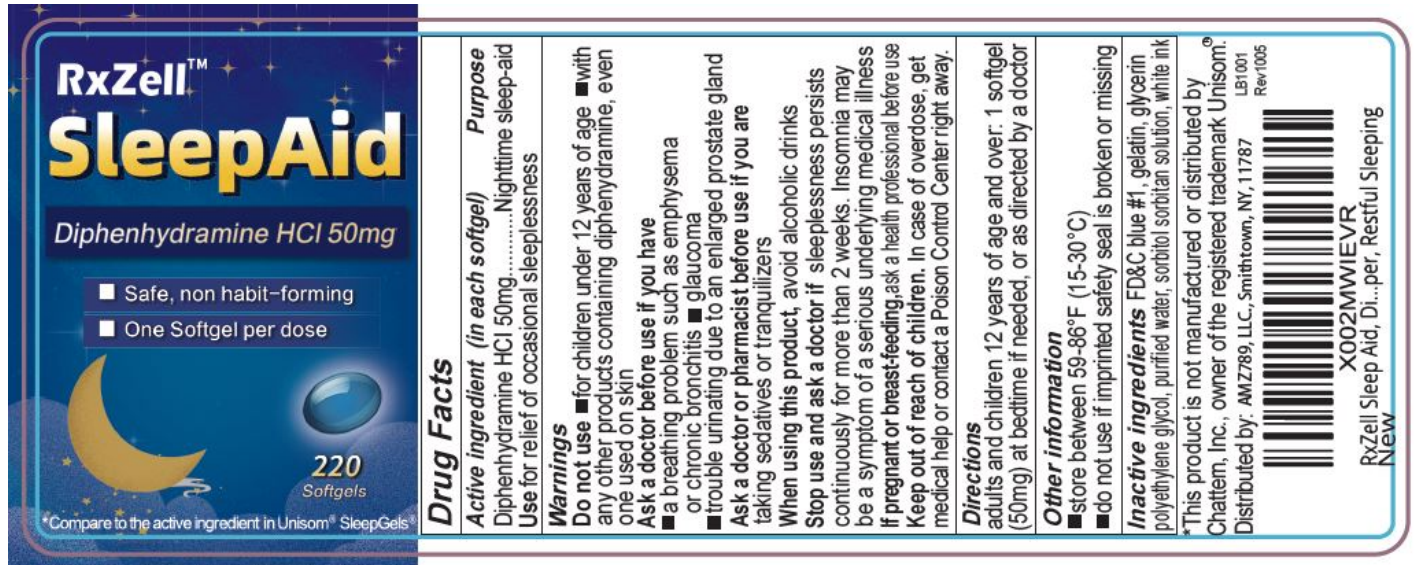
SleepAid

Diphenhydramine HCl 50mg

Non Habit-Forming

Easy to Swallow

Package Label Principal Display Panel



DIPHENHYDRAMINE HYDROCHLORIDE 50MG

diphenhydramine hydrochloride 50mg capsule, liquid filled

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73629-001
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
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
GLYCERIN (UNII: PDC6A3C0OX)	
GELATIN (UNII: 2G86QN327L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Product Characteristics

Color	blue (clear)	Score	no score
Shape	OVAL	Size	13mm
Flavor		Imprint Code	PC13
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73629-001-22	220 in 1 BOTTLE; Type 0: Not a Combination Product	09/28/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M010	09/21/2020	

Labeler - AMZ789 LLC (117410213)

Revised: 11/2024

AMZ789 LLC