# SOMNITABS- diphenhydramine hydrochloride tablet HARRIS TEETER

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.

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HTE - 1019 - 2019-1010

**Drug Facts** 

## Active ingredient (in each tablet)

Diphenhydramine HCl 25 mg

# Purpose

Nighttime sleep-aid

#### Use

helps to 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

## 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 drinks

**Stop use and ask a doctor if** sleeplessness persists continuously for more than 2 weeks. Insomnia may be a symptom of 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 right away.

#### **Directions**

 adults and children 12 years of age and over: take 2 tablets at bedtime if needed, or as directed by a doctor

### Other information

- each tablet contains: calcium 85 mg
- store at room temperature 15°-30°C (59°-86°F)

• retain carton for complete product information

# **Inactive ingredients**

colloidal silicon dioxide, croscarmellose sodium, dibasic calcium phosphate, FD&C blue #1, magnesium stearate, microcrystalline cellulose

## PRINCIPAL DISPLAY PANEL

Harris Teeter

Somnitabs

Compare to the Active Ingredient in Sominex

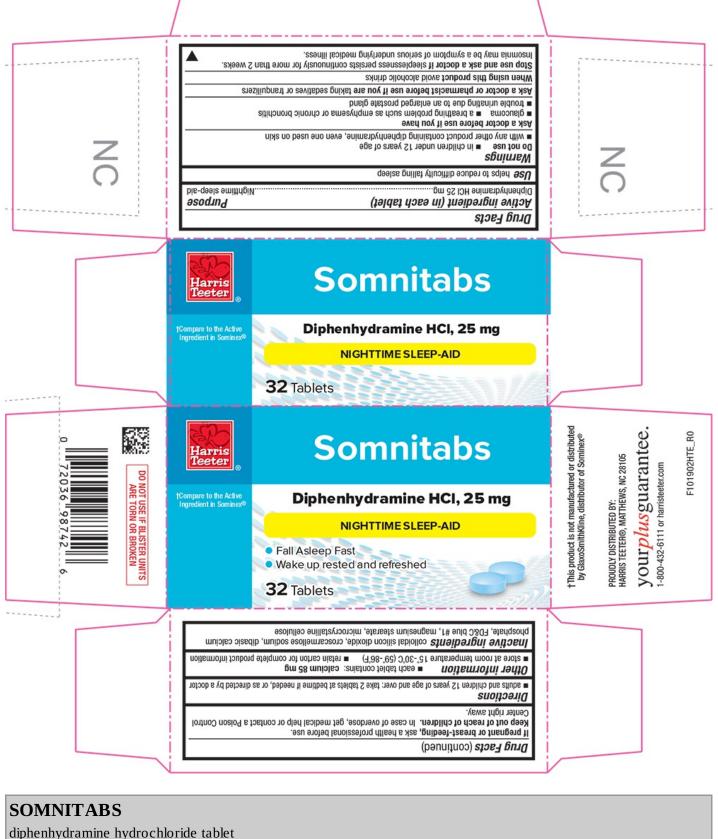
Diphenhydramine HCl, 25 mg

NIGHTTIME SLEEP-AID

Fall Asleep Fast

Wake up rested and refreshed

32 Tablets



Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72036-019
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
<b>DIPHENHYDRAMINE HYDRO CHLO RIDE</b> (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg

Inactive Ingredients		
Ingredient Name	Strength	
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)		
CROSCARMELLOSE SODIUM (UNII: M28 OL1HH48)		
CALCIUM PHO SPHATE, DIBASIC, ANHYDRO US (UNII: L11K75P92J)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
CELLULOSE, MICRO CRYSTALLINE (UNII: OP1R32D61U)		

Product Characteristics			
Color	blue	Score	no score
Shape	ROUND	Size	10 mm
Flavor		Imprint Code	93XF;57344
Contains			

Packaging			
# Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1 NDC:72036-019-02	4 in 1 CARTON	10/06/2011	
1	8 in 1 BLISTER PACK; 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	10/06/2011		

# Labeler - HARRIS TEETER (047279351)

Revised: 10/2019 HARRIS TEETER