NYTOL- diphenhydramine hcl capsule, gelatin coated
Medtech Products Inc.

Nytol Gelcap 50 mg_Export only 63029-928

Drug Facts

Active ingredient

Purpose

Diphenhydramine Hydrochloride 50 mg	Nighttime sleep aid

Uses/ Indications

- helps to reduce difficulty falling asleep
- relieves occasional sleeplessness

Warnings

Do not use

- if you are elderly, as this drug may cause excitation rather than sedation
- with any other product containing diphenhydramine, even one used on skin
- in children under 12 years of age

Ask a doctor or pharmacist before use if you

- have
  - a breathing problem such as emphysema or chronic bronchitis
  - glaucoma
- difficulty urinating
- take sedatives or tranquilizers
- are pregnant or breast-feeding

When using this product

avoid drinking alcohol

Stop use and ask a doctor if

sleeplessness continues for more than 2 weeks. Sleeplessness may be a symptom of a serious underlying medical illness.

Keep out of reach of children.

In case of overdose, call a poison control centre or get medical help right away.

Directions

Adults and children 12 years and over:

take 1 capsule at bedtime if needed, or as directed by a doctor

- if you feel drowsy in the morning, consult a doctor or pharmacist, as you may require a lower dosage (25 mg)
- do not take more than directed

Other information

Store between 15 & 30 °C

Inactive ingredients

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

Questions?

1-800-465-8811 www.nytol.ca

PRINCIPAL DISPLAY PANEL

Extra Strength Ultra-fort

Nytol®/MD

QUICKGELS®

GÉLULES-RAPIDES^{MC}

Extra Strength Ultra-fort

50 mg Diphenhydramine Hydrochloride Capsules USP

Capsules de chlorhydrate de diphenhydramine 50 mg USP

16 soft gels

gélules molles

DIN 02237169

NYTOL

diphenhydramine hcl capsule, gelatin coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63029-928
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)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
SORBITAN (UNII: 6O92ICV9RU)	
SORBITOL (UNII: 506T60A25R)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics			
Color	blue	**Score**	no score
Shape	OVAL	**Size**	1mm
Flavor		**Imprint Code**	N
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63029-928-16	2 in 1 CARTON	06/01/2012	
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
Export only		06/01/2012	

Labeler - Medtech Products Inc. (122715688)

Revised: 1/2024 Medtech Products Inc.