PAIN RELIEVER PM- acetaminophen and diphenhydramine hydrochloride tablet Health Pharma USA LLC

Pain Reliever PM - Acetaminophen and Diphenhydramine HCI

Drug Facts

Active ingredients (in each caplet)	Purpose
Acetaminophen 500 mg	Pain reliever
Diphenhydramine HCl 25 mg	Nighttime sleep aid

Uses

temporary relief of occasional headaches and minor aches and pains with accompanying sleeplessness

Warnings

Liver warning

This product contains acetaminophen. Severe liver damage may occur if you take

- more than 4,000 mg of acetaminophen in 24 hours
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

Do not use

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- with any product containing diphenhydramine, even one used on skin
- in children under 12 years of age
- if you have ever had an allergic reaction to this product or any of its ingredients

Ask a doctor before use if you have

- liver disease
- a breathing problem such as emphysema or chronic bronchitis
- trouble urinating due to an enlarged prostate gland
- glaucoma

Ask a doctor or pharmacist before use if you are

- taking the blood thinning drug warfarin
- taking sedatives or tranquilizers

When using this product

drowsiness will occur

- avoid alcoholic drinks
- do not drive a motor vehicle or operate machinery

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.
- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur

These could be signs of a serious condition.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children.

Overdose warning

Taking more than the recommended dose (overdose) may cause liver damage. In case of overdose, get medical help or contact a Poison Control Center right away. (1-800-222-1222) Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions

- do not take more than directed (see overdose warning)
- adults and children 12 years and over: take 2 caplets at bedtime
- do not take more than 2 caplets of this product in 24 hours
- **children under 12 years:** do not use this adult product in children under 12 years of age; this will provide more than the recommended dose (overdose) and may cause liver damage

Other information

- store between 20-25°C (68-77°F)
- do not use if pouch is torn or open
- see side panel for lot number and expiration date

Inactive ingredients

Acetaminophen

Diphenhydramine Hydrochloride

Microcrystalline cellulose

Pregelatinized starch

Stearic Acid

Colloidal Silicon Dioxide

Magnesium Stearate

Polyvinyl Alcohol

Polyethylene Glycol

PVP-K30

Sodium starch glycolate

Titanium Dioxide

FD&C blue no. 1

FD&C blue no. 2

Questions or comments?

Call toll free 1-844-832-1138 Monday through Friday 9AM -5PM EST or www.healthpharma.us

PRINCIPAL DISPLAY PANEL

See New Warnings Information & Directions

Compare to the Active Ingredients in

Tylenol PM ®*

PAIN RELIEVER PM

★Pain Reliever★Nighttime Sleep Aid

Acetaminophen, Diphenhydramine HCI



Health Pharma USA LLC

1600 Hart Street, Rahway, NJ 07065 Phone: 732-540-8421

FINISHED BULK LABEL of BOXES

PRODUCT NAME: Acetaminophen, Diphenhydramine HCI Film Coated Light Blue Caplets, 500 mg, 25 mg				
PRODUCT Code # PC143 Batch # Labeler code -71679-143-0				
NET WEIGHT (KG)		Caplets /BOX		
16.00 kg		25,000 Caplets		
		10		

BULK PACKAGING: 2 MIL clear Polyethylene Bag in Brown Flute Double Wall Corrugated Boxes.

BULK PRODUCT: Package immediately upon receipt.

WARNING: Keep out of Children. This bulk shipment is intended for further processing only. Repackage contents as soon as possible. This is a shipping container only and will not protect contents from damage by heat, cold and moisture indefinitely. Manufacturer will not accept responsibility for spoilage or violations of the food, Drug and cosmetic act due to prolonged storage in this container. Contents must be repackaged prior to resale and must be labeled in full compliance with the requirement of the Food, Drug and Cosmetic Act.

STORAGE: Store at a controlled room temperature 15-30°C (59-86° F)

PAIN RELIEVER PM

acetaminophen and diphenhydramine hydrochloride tablet

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71679-143	
Route of Administration	ORAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	500 mg	
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg	

Inactive Ingredients			
Ingredient Name	Strength		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)			
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)			

STARCH, PREGELATINIZED CORN (UNII: O8232NY3SJ)

POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)

STEARIC ACID (UNII: 4ELV7Z65AP)

TITANIUM DIOXIDE (UNII: 15FIX9V2JP)

SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)

POVIDONE (UNII: FZ989GH94E)

Product Characteristics					
Color blue Score no score					
Shape	CAPSULE	Size	18mm		
Flavor		Imprint Code	APM		
Contains					

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:71679-143- 00	1 in 1 BOX	02/01/2021		
1		25000 in 1 BAG; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	02/01/2016	

Labeler - Health Pharma USA LLC (080804485)

Establishment			
Name	Address	ID/FEI	Business Operations
Health Pharma USA LLC		080804485	manufacture(71679-143)

Revised: 10/2023 Health Pharma USA LLC