# PAIN RELIEVER PM- acetaminophen, diphenhydramine hydrochloride tablet Walmart Inc.

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### Walmart Acetaminophen 500 mg and Diphenhydramine HCl 25 mg Tablets

### **Drug Facts**

### Active ingredients (in each caplet)

Acetaminophen 500 mg

Diphenhydramine HCI 25 mg

### **Purpose**

Pain reliever

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

**Allergy alert:** acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash

If a skin reaction occurs, stop use and seek medical help right away.

#### 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 other 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:** 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

#### Other information

• store between 20-25°C (68-77°F)

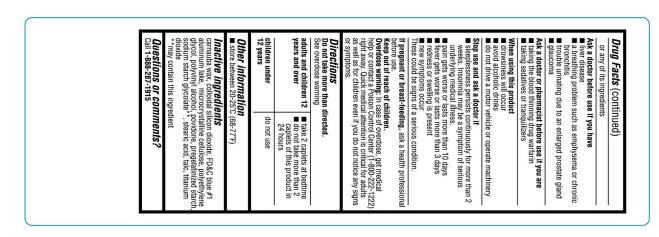
### Inactive ingredients

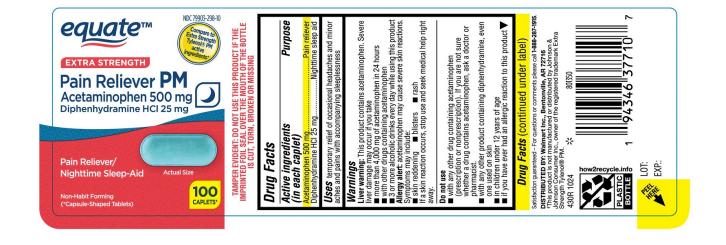
carnauba wax, colloidal silicon dioxide, FD&C blue#1 aluminum lake, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, pregelatinized starch, sodium starch glycolate\*\*, stearic acid, talc, titanium dioxide

\*\*may contain this ingredient

### Questions or comments?

Call 1-888-287-1915





# **PAIN RELIEVER PM**

acetaminophen, diphenhydramine hydrochloride tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79903-298
Route of Administration	ORAL		

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

Inactive Ingredients			
Ingredient Name	Strength		
STEARIC ACID (UNII: 4ELV7Z65AP)			
CARNAUBA WAX (UNII: R12CBM0EIZ)			
FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)			
TALC (UNII: 7SEV7J4R1U)			
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)			
POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E)			
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)			

Product Characteristics			
Color	blue	Score	no score
Shape	CAPSULE (Capsule shaped tablet)	Size	18mm
Flavor		Imprint Code	TCL430
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79903-298- 10	100 in 1 BOTTLE; Type 0: Not a Combination Product	01/15/2025	

Marketing Information			
Marketing Application Number or Monograph Category Citation		Marketing Start Marketing End Date Date	
OTC Monograph Drug	M013	01/15/2025	

# Labeler - Walmart Inc. (051957769)

# **Registrant -** TIME CAP LABORATORIES INC (037052099)

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
TIME CAP LABORATORIES INC		037052099	manufacture(79903-298)	

Revised: 12/2024 Walmart Inc.