# PAIN RELIEVER PM- acetaminophen, diphenhydramine hcl tablet, coated TARGET Corporation

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# **Drug Facts**

# Active ingredients (in each caplet)

# Acetaminophen 500 mg

Diphenhydramine HCl 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 products 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 a serious underlying medical illness.
- pain gets worse or last more than 10 days
- fever gets worse or last 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 (1-800-222-1222) right away. 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)
- avoid high humidity and excessive heat

# **Inactive ingredients**

croscarmellose sodium, FD&C blue #1 aluminum lake, FD&C blue #2 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone K30, pregelatanized starch, purified water, silicon dioxide,

## Questions or comments?

Call **1-800-910-6874** 

# **Principal Display Panel**

Compare to the active ingredients in Extra Strength Tylenol® PM\*

# **Extra Strength**

## Pain Reliever PM

Acetaminophen 500 mg/Pain Reliever

Diphenhydramine HCl 25 mg

Nighttime Sleep Aid

For adults

**CAPLETS** 

†This product is not manufactured or distributed by McNeil Consumer Healthcare, distributor of Extra Strength Tylenol® PM.

# TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

# KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

Distributed by

**Target Corporation** 

Minneapolis, MN 55403

# **Package Label**

Exp. Date: Lot No .: W 0 86

094 01 0372 R00 C-002262-01-034-0000

PLD-A596A FC008752

# Drug Facts (continued)

Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfarin taking sedatives or tranquilizers

# When using this product

microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone K30, pregelatinized starch, purified

water, silicon dioxide, sodium starch glycolate, talc,

titanium dioxide

FD&C blue #1 aluminum take, FD&C blue #2 aluminum

lake, hypromelioses, magnesium stearate,

Inactive ingredients croscarmellose sodium,

**Drug Facts** (continued

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

- Stop use and ask a doctor if
- 2 weeks. Insomnia may be a symptom of a serious sleeplessness persists continuously for more than
  - pain gets worse or lasts more than 10 days fever gets worse or lasts more than 3 days underlying medical illness.
    - redness or swelling is present

Love it or your money back.

Satisfaction guaranteed –

Questions or comments? Call 1-800-910-6874

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These could be signs of a serious condition.

# f pregnant or breast-feeding, ask a health professional

Distributed by Target Corporation Minneapolis, MN 55403

NDC 11673-754-10

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or distributed by McNeil Consume

Healthcare, distributor of Extra Strength Tylenol\* PM.

TM & @2024 Target Brands, Inc. This product is not manufactured

Product of India

# Directions

symptoms.

# do not take more than directed (see Overdose

- take 2 caplets at bedtime. Do not take more than adults and children 12 years of age and over.
  - children under 12 years of age; do not use caplets of this product in 24 hours.

# Other information

■ store between 20-25°C (68-77°F)
■ avoid high humidity and excessive heat

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION

# Orug Facts

# Active ingredients 'in each caplet

Purposes

Pain reliever Diphenhydramine HCl 25 mg.

. Nighttime sleep-aid Uses temporary relief of occasional headaches and

minor aches and pains with accompanying sleeplessness

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

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

Allergy alert: Acetaminophen may cause severe skin product

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- (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor with any other drug containing acetaminophen
  - 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
  - Ask a doctor before use if you have or any of its ingredients
- a breathing problem such as emphysema or chronic trouble urinating due to an enlarged prostate gland liver disease bronchitis

Actual Size

100 Caplets

glaucoma

Compare to active ingredients in Extra Strength Tylenol® PM\*

Strength a Reliever

Acetaminophen 500 mg / Pain Reliever Diphenhydramine HCl 25 mg Nighttime Sleep Aid For adults

100 CAPLETS

# TARGET Extra Strength Pain Reliever PM

# **PAIN RELIEVER PM**

acetaminophen, diphenhydramine hcl tablet, coated

<b>Product Information</b>			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-754
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	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)		
HYPROMELLOSES (UNII: 3NXW29V3WO)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)		
POVIDONE K30 (UNII: U725QWY32X)		
STARCH, CORN (UNII: O8232NY3SJ)		
WATER (UNII: 059QF0KO0R)		
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)		
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)		
TALC (UNII: 7SEV7J4R1U)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		
ALUMINUM OXIDE (UNII: LMI2606933)		

Product Characteristics				
Color	blue	Score	no score	
Shape	OVAL	Size	18mm	
Flavor		Imprint Code	P525	
Contains				

ı	Packaging			
#	# Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:11673- 754-10	1 in 1 BOX	08/01/2023	
1		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
	Marketing Information			
M	larketing	Information		
M	larketing Marketing Category	Information  Application Number or Monograph  Citation	Marketing Start Date	Marketing End Date

# Labeler - TARGET Corporation (006961700)

Revised: 10/2023 TARGET Corporation