PAIN RELIEVER PM EXTRA STRENGTH- acetaminophen, diphenhydramine hcl tablet, coated Walgreens

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.

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

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. 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 15-30°C (59-86°F)
- avoid high humidity and excessive heat

Inactive ingredients

carnauba wax*, croscarmellose sodium*, FD&C blue #1 aluminum lake, FD&C blue #2 aluminum lake, hypromellose, magnesium stearate*, microcrystalline cellulose, polyethylene glycol, polysorbate 80*, polyvinyl alcohol*, povidone K30, pregelatanized starch, purified water*, silicon dioxide*, sodium starch glycolate*, stearic acid*, talc*, titanium dioxide

*contains one or more of these ingredienrts

Questions or comments?

Call 1-877-753-3935 Monday-Friday 9AM-5PM EST

Principal Display Panel

Compare to Extra Strength Tylenol® PM active ingredients^{††}

Pain Reliever PM

ACETAMINOPHEN 500 mg/ PAIN RELIEVER

DIPHENHYDRAMINE HCl 25 mg/ NIGHTTIME SLEEP AID

NIGHTTIME EXTRA STRENGTH CAPLETS

CAPLETS**

(**CAPSULES-SHAPED TABLETS)

††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: WALGREEN CO.

200 WILMOT RD., DEERFIELD, IL 60015

walgreens.com

Package Label

Exp. Date

ORG0718-F

FC005062 PLD-D134H

ITEM 283596 W10024-0718-L 19 5965 N

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DISTRIBUTED BY: WALGREEN CO. 200 WILMOT RD., DEERFIELD, IL 60015 **00%** SATISFACTION

Orug Facts (continued)

FD&C blue #2 aluminum lake, hypromellose, magnesium stearate*, microcrystalline cellulose, polyethylene glycol croscarmellose sodium*, FD&C blue #1 aluminum lake, sodium starch glycolate", stearic acid", talc", trtanium polysorbate 80°, polyvinyl alcohol", povidone K30, pregelatinized starch, purified water", silicon dioxide* inactive ingredients camauba wax

contains one or more of these ingredients

Call 1-877-753-3935 Monday-Friday 9AM-5PM EST Questions or comments?

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



Drug Facts (continued)

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

Purposes

Active ingredients

Orug Facts

in each caplet,

taking sedatives or tranquilizers

When using this product

Nighttime sleep-aid

Diphenhydramine HCI 25 mg.

 avoid alcoholic drinks drowsiness will occur

do not drive a motor vehicle or operate machinery

 sleeplessness persists continuously for more than Stop use and ask a doctor if

2 weeks. Insomnia may be a symptom of a serious

underlying medical illness. new symptoms occur

redness or swelling is present

pain gets worse or lasts more than 10 days

 fever gets worse or lasts more than 3 days These could be signs of a serious condition If pregnant or breast-feeding, ask a health professional

before use

fa skin reaction occurs, stop use and seek medical help

■ skin reddening ■ blisters ■ rash

reactions. Symptoms may include:

(prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor

with any other drug containing acetaminophen

Do not use

right away.

with any other product containing diphenhydramine

or pharmacist

even one used on skin

Overdose warming: 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 Keep out of reach of children. symptoms

Directions

 do not take more than directed (see Overdose warning) adults and children 12 years of age and over

take 2 caplets at bedtime. Do not take more than caplets of this product in 24 hours.

■ store between 15-30°C (59-86°F)
■ avoid high humidity and excessive heat

children under 12 years of age: do not use

Other information

a breathing problem such as emphysema or chronic

Ask a doctor before use If you have

liver disease

bronchitis glaucoma

or any of its ingredients

trouble urinating due to an enlarged prostate gland

product

3 or more alcoholic drinks every day while using this

 with other drugs containing acetaminophen Severe liver damage may occur if you take:

Allergy alert: Acetaminophen may cause severe skin

Walgreens

Uses :

minor aches and pains with accompanying sleeplessness.

iver warning: This product contains acetaminophen. ■ more than 4,000 mg of acetaminophen in 24 hours

Narnings

temporary relief of occasional headaches and

Compare to Extra Strength Tylenol® PM active ingredients#

in children under 12 years of ageif you have ever had an allergic reaction to this product

NDC 0363-4470-10

ACETAMINOPHEN 500 mg / PAIN RELIEVER DIPHENHYDRAMINE HCI 25 mg / NIGHTTIME SLEEP AID

NIGHTTIME

EXTRA STRENGTH

CAPLETS

ACTUAL SIZE

CAPLETS**

(**CAPSULE-SHAPED TABLETS)

PAIN RELIEVER PM EXTRA STRENGTH

acetaminophen, diphenhydramine hcl tablet, coated

Product Type HUMAN OTC DRUG Item Cod	e (Source)	NDC:0363-4470
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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
CARNAUBA WAX (UNII: R12CBM0EIZ)	
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)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
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)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	blue	Score	no score
Shape	CAPSULE	Size	18mm
Flavor		Imprint Code	S525;P525;G651
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363- 4470-10	1 in 1 BOX	12/30/2014	11/30/2025
1		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:0363- 4470-15	1 in 1 BOX	12/30/2014	11/30/2025
2		150 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:0363- 4470-52	250 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/30/2014	11/30/2025
4	NDC:0363- 4470-11	1 in 1 BOX	12/30/2014	11/30/2025
4		110 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
5	NDC:0363- 4470-30	1 in 1 BOX	12/30/2014	11/30/2025
5		30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
6	NDC:0363- 4470-24	1 in 1 BOX	12/30/2014	11/30/2025
6		24 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	12/30/2014	11/30/2025

Labeler - Walgreens (008965063)

Revised: 11/2022 Walgreens