

**ACETAMINOPHEN AND DIPHENHYDRAMINE HCL- acetaminophen and
diphenhydramine hcl tablet
PUBLIX SUPERMARKETS, INC**

**Extra Strength
Pain Relief PM**

ACETAMINOPHEN, 500 mg DIPHENHYDRAMINE HCl, 25 mg

- **Pain reliever/nighttime sleep-aid**
- **Non-habit forming**
- **Contains no aspirin**

Active ingredients (in each caplet)

Acetaminophen USP, 500 mg

Diphenhydramine HCl USP, 25 mg

Purpose

Pain reliever

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

- 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

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). See USP Controlled Room Temperature.
- see end panel for lot number and expiration date

Inactive ingredients

carnauba wax, colloidal silicon dioxide, croscarmellose sodium, FD&C blue #1 aluminum lake, FD&C blue #2 aluminum lake, hypromellose, microcrystalline cellulose, polyethylene glycol, polysorbate 80, povidone k-30, pregelatinized starch, stearic acid, titanium dioxide

Questions or comments?

call **1-877-770-3183** Mon-Fri 8:00 AM EST to 5:00 PM PST

Principal display panel

COATING
FREE AREA

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Drug Facts (continued)

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Drug Facts (continued)

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Inactive ingredients

croscarmellose sodium, FD&C blue #1 aluminum lake, FD&C blue #2 aluminum lake, hypromellose, microcrystalline cellulose, polyethylene glycol, polyorbate 80, povidone K-30, pregelatinized starch, stearic acid, titanium dioxide

Questions or comments?

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*This product is not manufactured or distributed by Kenvue Inc., owner of the registered trademark Extra Strength Tylenol® PM Caplets.

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PUBLIX GUARANTEE: COMPLETE SATISFACTION OR YOUR MONEY BACK



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Pain
Relief PM**

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 DIPHENHYDRAMINE HCl, 25 mg

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50 Caplets*
 *Caplet-enclosed tablets

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

ACTUAL SIZE



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

READ AND KEEP CARTON FOR COMPLETE WARNINGS AND INFORMATION

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

Drug Facts

Active ingredients (in each caplet) Purpose

Acetaminophen USP, 500 mg Pain reliever
 Diphenhydramine HCl USP, 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

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- with other drugs containing acetaminophen
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Allergy alert: acetaminophen may cause severe skin reactions. Symptoms may include:

- skin redness
- hives
- rash
- if a skin reaction occurs, stop use and seek medical help right away

Do not use

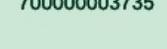
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Publix.

Lot
Exp.

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NDC 41415-661-05



ACETAMINOPHEN AND DIPHENHYDRAMINE HCL

acetaminophen and diphenhydramine hcl tablet

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:41415-661 |
| Route of Administration | ORAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|-------------------|----------|
| DIPHENHYDRAMINE (UNII: 8GTS82S83M) (DIPHENHYDRAMINE - UNII:8GTS82S83M) | DIPHENHYDRAMINE | 25 mg |
| ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D) | ACETAMINOPHEN | 500 mg |

Inactive Ingredients

| Ingredient Name | Strength |
|--|----------|
| CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U) | |
| CARNAUBA WAX (UNII: R12CBM0EIZ) | |
| TITANIUM DIOXIDE (UNII: 15FIX9V2JP) | |
| CROSCARMELOSE SODIUM (UNII: M28OL1HH48) | |
| POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ) | |
| POLYSORBATE 80 (UNII: 6OZP39ZG8H) | |
| SILICON DIOXIDE (UNII: ETJ7Z6XBU4) | |
| STARCH, CORN (UNII: O8232NY3SJ) | |
| FD&C BLUE NO. 2 (UNII: L06K8R7DQK) | |
| STEARIC ACID (UNII: 4ELV7Z65AP) | |
| POVIDONE K30 (UNII: U725QWY32X) | |
| HYPROMELLOSES (UNII: 3NXW29V3WO) | |
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD) | |

Product Characteristics

| | | | |
|----------|------|--------------|----------|
| Color | blue | Score | no score |
| Shape | OVAL | Size | 17mm |
| Flavor | | Imprint Code | G651 |
| Contains | | | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:41415-661-05 | 50 in 1 BOTTLE; Type 0: Not a Combination Product | 08/02/2024 | |
| 2 | NDC:41415-661-10 | 100 in 1 BOTTLE; Type 0: Not a Combination Product | 08/02/2024 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | M013 | 08/02/2024 | |

Labeler - PUBLIX SUPERMARKETS, INC (006922009)

Revised: 12/2025

PUBLIX SUPERMARKETS, INC