

**ACETAMINOPHEN AND DIPHENHYDRAMINE HYDROCHLORIDE - acetaminophen and diphenhydramine hydrochloride tablet
WALMART INC.**

Drug Facts

Active ingredients (in each gelcap)

Acetaminophen USP 500 mg
Diphenhydramine hydrochloride USP 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:

- with other drugs containing acetaminophen
- more than 4,000 mg of acetaminophen in 24 hours
- 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
- if you are allergic to acetaminophen or any of the inactive ingredients in this product
- in children under 12 years of age

Ask a doctor before use if you have

- liver disease.
- a breathing problem such as emphysema or chronic bronchitis
- difficulty in urination due to enlargement of the 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 beverages
- do not drive a motor vehicle or operate machinery

Stop use and ask a doctor if

- pain gets worse or lasts more than 10 days
- 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
- 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.

Keep out of reach of children.

In case of accidental overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away. Prompt 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**
- adults and children 12 years and over
 - take 2 gelcaps at bedtime
 - do not take more than 2 gelcaps of this product in 24 hours.
- children under 12 years: do not use

Other information

- avoid high humidity

- store at 20° to 25°C (68° to 77°F)
- use by expiration date on package

Inactive ingredients

ammonium hydroxide, black iron oxide, colloidal silicon dioxide, croscarmellose sodium, FD&C blue #1, FD&C red #3, gelatin, hydroxypropyl cellulose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, povidone, pregelatinized starch (maize), propylene glycol, shellac glaze, talc, and titanium dioxide.

Questions or comments?

call 1-888-287-1915 (Monday-Friday 8:30 AM to 5:00 PM EST)

**DISTRIBUTED BY: Walmart Inc.,
Bentonville, AR 72716**

PRODUCT OF INDIA

**PACKAGE LABEL-PRINCIPAL DISPLAY PANEL 500 mg/25 mg (80 Gelcaps
Bottle)**

equate™

NDC 79903-267-80

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

EXTRA STRENGTH

**Pain Reliever PM
Acetaminophen 500 mg
Diphenhydramine HCl 25 mg**

**Pain Reliever/Nighttime Sleep-Aid
Non-habit forming**

Actual Size

**80
GELCAPS**

equate™

NDC 79905-267-80



EXTRA STRENGTH

Pain Reliever PM
Acetaminophen 500 mg
Diphenhydramine HCl 25 mg



Pain Reliever/Nighttime Sleep-Aid
Non-habit forming



Actual Size

80
GELCAPS

Do not use if printed foil seal under cap is torn or missing.

Drug Facts

Active ingredients (in each gelcap)	Purpose
Acetaminophen USP 500 mg	Pain reliever
Diphenhydramine hydrochloride 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:

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

Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include:

DISTRIBUTED BY: Walmart Inc.,
Bentonville, AR 72716
PRODUCT OF INDIA
*This product is not manufactured or distributed by
McNeil Consumer Healthcare, owner of the
registered trademark Extra Strength Tylenol® PM.

Satisfaction guaranteed - For questions
or comments please call 1-888-287-1915.



P1434530 LM-5847

Unvarnished Zone
(dotted line not for printing)
12 x 27 mm

Lift Here

Drug Facts (continued)

■ skin redness ■ 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
- if you are allergic to acetaminophen or any of the inactive ingredients in this product
- in children under 12 years of age

Ask a doctor before use if you have

- liver disease.
- a breathing problem such as emphysema or chronic bronchitis
- difficulty in urination due to enlargement of the 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 beverages
- do not drive a motor vehicle or operate machinery

Stop use and ask a doctor if

- pain gets worse or lasts more than 10 days
- 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
- 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.

Keep out of reach of children. In case of accidental overdose, get medical help or contact a Poison Control Center

HINGE

HINGE

Drug Facts (continued)

(1-800-222-1222) right away. Prompt 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
- adults and children 12 years and over
- take 2 gelcaps at bedtime
- do not take more than 2 gelcaps of this product in 24 hours.
- children under 12 years: do not use

Other information

- avoid high humidity
- store at 20° to 25°C (68° to 77°F)
- use by expiration date on package

Inactive ingredients

ammonium hydroxide, black iron oxide, colloidal silicon dioxide, croscarmellose sodium, FD&C blue #1, FD&C red #3, gelatin, hydroxypropyl cellulose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, povidone, pregelatinized starch (maize), propylene glycol, shellac glaze, talc, and titanium dioxide.

Questions or comments? call 1-888-287-1915 (Monday-Friday 8:30 AM to 5:00 PM EST)

P1434530

LM-5847



ACETAMINOPHEN AND DIPHENHYDRAMINE HYDROCHLORIDE

acetaminophen and diphenhydramine hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79903-267
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg

Inactive Ingredients

Ingredient Name	Strength
AMMONIA (UNII: 5138Q19F1X)	
FERROSFERRIC OXIDE (UNII: XM0M87F357)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 3 (UNII: PN2ZH5LOQY)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
HYDROXYPROPYL CELLULOSE (110000 WAMW) (UNII: 5Y0974F5PW)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE 101 (UNII: 7T9FYH5QMK)	
MICROCRYSTALLINE CELLULOSE 102 (UNII: PNR0YF693Y)	
POLYETHYLENE GLYCOL 4000 (UNII: 4R4HFI6D95)	
POVIDONE K90 (UNII: RDH86HJV5Z)	
STARCH, CORN (UNII: O8232NY3SJ)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SHELLAC (UNII: 46N107B71O)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	BLUE (Dark blue and Light blue with white band)	Score	no score
Shape	CAPSULE (Biconvex)	Size	20mm
Flavor		Imprint Code	T;6
Contains			

Packaging

#	Item Code	Package Description	Marketing Start	Marketing End
---	-----------	---------------------	-----------------	---------------

#	Item Code	Package Description	Date	Date
1	NDC:79903-267-80	80 in 1 BOTTLE; Type 0: Not a Combination Product	06/28/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	06/28/2024	

Labeler - WALMART INC. (051957769)

Registrant - Aurohealth LLC (078728447)

Establishment			
Name	Address	ID/FEI	Business Operations
APL HEALTHCARE LIMITED		650844777	ANALYSIS(79903-267) , MANUFACTURE(79903-267)

Revised: 7/2024

WALMART INC.