

**PAIN RELIEVER PM- acetaminophen, diphenhydramine hcl tablet, film coated  
WALMART INC.**

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**Equate 44-235**

***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 product containing diphenhydramine, even one used on skin
- in children under 12 years of age
- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- if you have ever had an allergic reaction to this product or any of its ingredients

**Ask a doctor before use if you have**

- a breathing problem such as emphysema or chronic bronchitis
- liver disease

- glaucoma
- difficulty in urination due to enlargement of the prostate gland

**Ask a doctor or pharmacist before use if you are**

- taking the blood thinning drug warfarin
- taking sedatives or tranquilizers

**When using this product**

- avoid alcoholic beverages
- drowsiness will occur
- 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.
- redness or swelling is present
- new symptoms occur
- 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.

**Keep out of reach of children.**

In case of accidental overdose, get medical help or contact a Poison Control Center 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 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 at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- see end flap for expiration date and lot number

***Inactive ingredients***

colloidal silicon dioxide, corn starch, croscarmellose sodium, FD&C blue #1 aluminum lake, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, stearic acid, talc, titanium dioxide

**Questions or comments?**

**1-888-287-1915**

**Principal Display Panel**

**equate™**

NDC 79903-307-10

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  
For Ages 12 Years and Over

Actual Size

**100  
CAPLETS**

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

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

50844 REV0521A23512

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

Satisfaction guaranteed - Or we'll replace it or give you your money  
back. For questions or comments please call **1-888-287-1915**.

**equate™** NDC 79903-307-10

**EXTRA STRENGTH**

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

Compare to Extra Strength Tylenol's PM active ingredients\*

Pain Reliever  
 Nighttime Sleep-Aid

Actual Size

Non-Habit Forming  
 For Ages 12 Years and Over

100 CAPLETS

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

**Drug Facts**

**Active ingredients (in each caplet)** Purpose  
 Acetaminophen 500 mg ..... Pain reliever  
 Diphenhydramine HCl 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  
 ■ 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 product containing diphenhydramine, even one used on skin ■ in children under 12 years of age

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

**DISTRIBUTED BY:** Walmark Inc., Bentonville, AR 72716  
 \*This product is not manufactured or distributed by Wm. W. Green, owner of the registered trademark Extra Strength Tylenol® PM.

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 PLASTIC BOTTLE

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PEEL HERE FOR MORE DRUG FACTS

**Drug Facts (continued)**

■ with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.  
 ■ 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 ■ glaucoma  
 ■ a breathing problem such as emphysema or chronic bronchitis  
 ■ difficulty in urination due to enlargement of the prostate gland

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

**When using this product**  
 ■ avoid alcoholic beverages  
 ■ do not drive a motor vehicle or operate machinery  
 ■ drowsiness will occur

**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 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.**  
 Keep out of reach of children. In case of accidental overdose, get medical help or contact a Poison Control Center 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 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 at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)  
 ■ use by expiration date on package

**Inactive ingredients** colloidal silicon dioxide, corn starch, croscarmellose sodium, FD&C blue #1 aluminum lake, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, stearic acid, talc, titanium dioxide

**Questions or comments? 1-888-287-1915**

STOP PEELING

**Equate 44-235**

**PAIN RELIEVER PM**

acetaminophen, diphenhydramine hcl tablet, film coated

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:79903-307
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

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

**Inactive Ingredients**

Ingredient Name	Strength
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<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>CROSCARMELOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>FD&amp;C BLUE NO. 1 ALUMINUM LAKE</b> (UNII: J9EQA3S2JM)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>POLYVINYL ALCOHOL, UNSPECIFIED</b> (UNII: 532B59J990)	
<b>POVIDONE, UNSPECIFIED</b> (UNII: FZ989GH94E)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	

Product Characteristics			
<b>Color</b>	blue	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	17mm
<b>Flavor</b>		<b>Imprint Code</b>	44;235
<b>Contains</b>			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79903-307-10	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/23/2025	
2	NDC:79903-307-24	1 in 1 CARTON	11/25/2025	
2		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 Drug	M013	01/23/2025	

**Labeler** - WALMART INC. (051957769)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(79903-307)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(79903-307) , pack(79903-307)

**Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		832867894	manufacture(79903-307)

## **Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		868734088	manufacture(79903-307)

## **Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		967626305	pack(79903-307)

Revised: 11/2025

WALMART INC.