

**PMS RELIEF MAXIMUM STRENGTH- acetaminophen, pamabrom, pyrilamine maleate tablet, film coated**

**CVS Pharmacy**

*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.*

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**CVS 44-679**

***Active ingredients (in each caplet)***

Acetaminophen 500 mg

Pamabrom 25 mg

Pyrilamine maleate 15 mg

***Purpose***

Pain reliever

Diuretic

Antihistamine

***Uses***

for the temporary relief of these symptoms associated with menstrual periods:

- headache
- bloating
- cramps
- backache
- muscular aches
- irritability
- water-weight gain

***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.
- if you have ever had an allergic reaction to this product or any of its ingredients

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

- difficulty in urination due to enlargement of the prostate gland
- a breathing problem such as emphysema or chronic bronchitis
- liver disease
- glaucoma

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

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

**When using this product**

- drowsiness may occur
- avoid alcoholic beverages
- excitability may occur, especially in children
- alcohol, sedatives and tranquilizers may increase drowsiness
- use caution when driving a motor vehicle or operating machinery

**Stop use and ask a doctor if**

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

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 the recommended dose**
- adults and children 12 years and over:
  - take 2 caplets with water every 6 hours as needed
  - do not exceed 8 caplets in a 24 hour period or as directed by a doctor
- children under 12 years: ask a doctor

***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***

corn starch, croscarmellose sodium, crospovidone, FD&C blue #2 aluminum lake, FD&C red #40 aluminum lake, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, shellac wax, silicon dioxide, stearic acid, talc, titanium dioxide

***Questions or comments?***

**1-800-426-9391**

***Principal display panel***

CVSHealth™

Compare to the active ingredients in Maximum Strength Pamprin® Multi-Symptom\*

MAXIMUM STRENGTH

**PMS Relief**

**ACETAMINOPHEN** 500 mg

**PAMABROM** 25 mg, **PYRILAMINE MALEATE** 15 mg

Pain reliever, Diuretic, Antihistamine

Multi-symptom

Relieves the symptoms of premenstrual syndrome: cramps, bloating, headache, backache and water-weight gain

**24 COATED CAPLETS**

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

\*This product is not manufactured or distributed by Chattem, Inc., owner of the registered trademark Maximum Strength Pamprin® Multi-Symptom.

50844 REV0718B67908

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 registered trademark Maximum Strength Pamprin®  
 Focus Consumer Healthcare, LLC, owner of the

B-0231-679-08-H  
REV0718867908

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 V-11112

**Drug Facts (continued)**  
 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.

**Drug Facts (continued)**  
 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**  
 ■ difficulty in urination due to enlargement of the prostate gland  
 ■ a breathing problem such as emphysema or chronic bronchitis  
 ■ liver disease ■ glaucoma  
**Ask a doctor or pharmacist before use if you are**  
 ■ taking sedatives or tranquilizers  
 ■ taking the blood thinning drug warfarin  
**When using this product**  
 ■ drowsiness may occur  
 ■ avoid alcoholic beverages  
 ■ excitability may occur, especially in children  
 ■ alcohol, sedatives and tranquilizers may increase drowsiness  
 ■ use caution when driving a motor vehicle or operating machinery  
**Stop use and ask a doctor if**  
 ■ pain gets worse or lasts more than 10 days  
 ■ fever gets worse or lasts more than 3 days  
 ■ new symptoms occur  
 ■ redness or swelling is present  
 These could be signs of a serious condition.  
**If pregnant or breast-feeding, ask a health professional before use.**

**Drug Facts (continued)**  
**Directions**  
 ■ do not take more than directed  
 ■ adults and children 12 years and over:  
 ■ take 2 caplets with water every 6 hours as needed  
 ■ do not exceed 6 caplets in a 24 hour period or as directed by a doctor  
 ■ children under 12 years: ask a doctor

**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** corn starch, croscarmellose sodium, crospovidone, FD&C blue #2 aluminum lake, FD&C red #40 aluminum lake, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, stearic acid, silicon dioxide, stearic acid, talc, titanium dioxide

**Questions or comments?**  
 1-800-426-9391

**Drug Facts**  
**Active ingredients (in each caplet)**  
 Acetaminophen 500 mg, Pain reliever  
 Pamabrom 25 mg, Diuretic  
 Pyrilamine maleate 15 mg, Antihistamine

**Purpose**  
 Pain reliever  
 Diuretic  
 Antihistamine

**Uses** for the temporary relief of these symptoms associated with menstrual periods:  
 ■ headache ■ bloating ■ cramps  
 ■ backache ■ muscular aches  
 ■ irritability ■ water-weight gain

**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

**CVSHealth**

Compare to the active ingredients in  
 Maximum Strength Pamprin® Multi-Symptom\*

**MAXIMUM STRENGTH**  
**PMS Relief**  
**ACETAMINOPHEN 500 mg**  
**PAMABROM 25 mg, PYRILAMINE MALEATE 15 mg**  
 Pain reliever, Diuretic, Antihistamine  
 Multi-symptom  
 Relieves the symptoms of premenstrual syndrome: cramps, bloating, headache, backache and water-weight gain  
**24 COATED CAPLETS**



Actual Size

44-679

## PMS RELIEF MAXIMUM STRENGTH

acetaminophen, pamabrom, pyrilamine maleate tablet, film coated

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59779-879
Route of Administration	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg
PAMABROM (UNII: UA8U0KJM72) (BROMOTHEOPHYLLINE - UNII:FZG87K1MQ6)	PAMABROM	25 mg
PYRILAMINE MALEATE (UNII: R35D29L3ZA) (PYRILAMINE - UNII:HPE317O9TL)	PYRILAMINE MALEATE	15 mg

**Inactive Ingredients**

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
MAGNESIUM STEARATE (UNII: 70097M6B30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
CROSPVIDONE (UNII: 2S7830E561)	
SHELLAC (UNII: 46N107B71O)	

**Product Characteristics**

Color	PURPLE	Score	no score
Shape	OVAL	Size	17mm
Flavor		Imprint Code	44;679
Contains			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59779-879-27	1 in 1 CARTON	01/13/2015	
1		32 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:59779-879-08	1 in 1 CARTON	01/13/2015	
2		24 in 1 BOTTLE; Type 0: Not a Combination Product		

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part343	01/13/2015	

**Labeler** - CVS Pharmacy (062312574)

<b>Establishment</b>			
<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		038154464	PACK(59779-879)

<b>Establishment</b>			
<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		832867894	MANUFACTURE(59779-879)

<b>Establishment</b>			
<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		967626305	PACK(59779-879)

<b>Establishment</b>			
<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		832867837	PACK(59779-879)

<b>Establishment</b>			
<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		868734088	PACK(59779-879)

Revised: 5/2020

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