MENSTRUAL PAIN RELIEF- acetaminophen, pamabrom, pyrilamine maleate tablet, film coated Chain Drug Marketing Association, Inc.

Quality Choice 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

- if you have ever had an allergic reaction to this product or any of its ingredients
- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.

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 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, shellac wax, silicon dioxide, stearic acid, talc, titanium dioxide

Questions or comments?

1-800-426-9391

Principal Display Panel

QC QUALITY® CHOICE

NDC 63868-961-20

Compare to the Active Ingredients in Maximum Strength Pamprin® Multi-Symptom*

Maximum Strength

Menstrual Pain Relief

Multi-Symptom Acetaminophen - Pain Reliever

Pamabrom - Diuretic Pyrilamine Maleate - Antihistamine

Relieves: Cramps, Irritability, Headache, Bloating, Backache

20 Caplets

actual size

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

*This product is not manufactured or distributed by Focus Consumer Healthcare, LLC, owner of the registered trademark Maximum Strength Pamprin® Multi-Symptom. 50844 REV0718B67909

QC 100% SATISFACTION GUARANTEED Distributed by CDMA, Inc. Novi, MI 48375 www.qualitychoice.com

Www.qualitycnoice.com Questions: 800-935-2362



MENSTRUAL PAIN RELIEF

acetaminophen, pamabrom, pyrilamine maleate tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83324-057
Route of Administration	ORAL		

Active Ingredient/Active Moiety					
Ingredient Name	Basis of Strength	Strength			
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	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)	
CROSPOVIDONE, UNSPECIFIED (UNII: 2S7830E561)	
FD&C BLUE NO. 2 ALUMINUM LAKE (UNII: 4AQJ3LG584)	
FD&C RED NO. 40 ALUMINUM LAKE (UNII: 6T47AS764T)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E)	
SHELLAC (UNII: 46N107B710)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

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

I	Packaging						
7	# Item Code	Package Description	Marketing Start Date	Marketing End Date			
:	NDC:83324- 057-20	1 in 1 CARTON	06/21/2024				
:	L	20 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product					

Marketing Information				
Marketing	Application Number or Monograph	Marketing Start	Marketing End	

Category	Citation	Date	Date
OTC Monograph Drug	M013	06/21/2024	

Labeler - Chain Drug Marketing Association, Inc. (011920774)

Establishment					
Name	Address	ID/FEI	Business Operations		
LNK International, Inc.		038154464	pack(83324-057)		

Establishment				
Name	Address	ID/FEI	Business Operations	
LNK International, Inc.		832867837	manufacture(83324-057)	

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(83324-057)

Establishment					
Name	Address	ID/FEI	Business Operations		
LNK International, Inc.		967626305	pack(83324-057)		

Establishment					
Name	Address	ID/FEI	Business Operations		
LNK International, Inc.		117025878	manufacture(83324-057)		

Revised: 9/2025 Chain Drug Marketing Association, Inc.