MENSTRUAL COMPLETE- acetaminophen, caffeine, pyrilamine maleate tablet, film coated

Chain Drug Marketing Association, Inc.

Quality Choice 44-390

Active ingredients (in each caplet)

Acetaminophen 500 mg Caffeine 60 mg Pyrilamine maleate 15 mg

Purpose

Pain reliever Diuretic Antihistamine

Uses

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

- bloating
- headache
- muscle aches
- cramps
- backache
- fatique
- 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
- liver disease
- glaucoma
- a breathing problem such as emphysema or chronic bronchitis

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 may occur
- alcohol, sedatives, and tranquilizers may increase drowsiness
- avoid alcoholic beverages
- excitability may occur, especially in children
- use caution when driving a motor vehicle or operating machinery
- limit the use of caffeine-containing medications, foods, or beverages while taking this product because too much caffeine may cause nervousness, irritability, sleeplessness, and, occasionally, rapid heart rate. The recommended dose of this product contains about as much caffeine as a cup of coffee.

Stop use and ask a doctor if

- 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 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
 - repeat every 6 hours, as needed

- do not exceed 6 caplets per day
- children under 12 years: ask a doctor

Other information

- see end flap for expiration date and lot number
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)

Inactive ingredients

corn starch, croscarmellose sodium, crospovidone, hypromellose, magnesium stearate, microcrystalline cellulose, polydextrose, polyethylene glycol, povidone, silicon dioxide, stearic acid, titanium dioxide, triacetin

Questions or comments?

1-800-426-9391

Principal Display Panel

NDC 83324-017-24

QC_® QUALITY CHOICE

*Compare to the active ingredients in Midol® Complete

Menstrual Complete Acetaminophen 500 mg, Caffeine 60 mg

Pyrilamine maleate 15 mg

Pain Reliever / Diuretic / Antihistamine

Multi-Symptom Relief

24 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 Bayer HealthCare LLC, owner of the registered trademark Midol® Complete. 50844 REV0623B39008

Distributed by CDMA, Inc. Novi, MI 48375 www.qualitychoice.com Questions: 800-935-2362 SATISFACTION 100% QC GUARANTEED

MENSTRUAL COMPLETE

acetaminophen, caffeine, pyrilamine maleate tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83324-017
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	500 mg
CAFFEINE (UNII: 3G6A5W338E) (CAFFEINE - UNII:3G6A5W338E)	CAFFEINE	60 mg
PYRILAMINE MALEATE (UNII: R35D29L3ZA) (PYRILAMINE - UNII:HPE317O9TL)	PYRILAMINE MALEATE	15 mg

Inactive Ingredients	
Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)	
CROSPOVIDONE, UNSPECIFIED (UNII: 2S7830E561)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	

Product Characteristics				
Color	white	Score	no score	
Shape	OVAL	Size	17mm	
Flavor		Imprint Code	44;390	
Contains				

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
#	Item Code	Package Description		

1 NDC:83324- 017-24	1 in 1 CARTON	11/02/2023	
	24 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information					
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
M013	11/02/2023				
	Application Number or Monograph Citation	Application Number or Monograph Marketing Start Citation Date			

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

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(83324-017)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(83324-017) , pack(83324-017)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(83324-017)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(83324-017)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(83324-017)

Revised: 11/2023 Chain Drug Marketing Association, Inc.