MENSTRUAL COMPLETE MAXIMUM STRENGTH- acetaminophen, caffeine and pyrilamine maleate tablet, film coated
L.N.K. International, Inc.

Quality Plus 44-390 Maximum Strength

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
- water-weight gain
- cramps
- backache
- fatigue
- muscle aches

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
- a breathing problem such as emphysema or chronic bronchitis
- 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
- avoid alcoholic beverages
- excitability may occur, especially in children
- be careful when driving a motor vehicle or operating machinery
- alcohol, sedatives, and tranquilizers may increase drowsiness
- you may get drowsy
- 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 heartbeat. 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 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
- 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 25ºC (77ºF); excursions permitted between 15º-30ºC (59º-86ºF)
- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN**

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

QUALITY
PLUS

NDC 50844-390-21

*Compare to the active ingredients in Midol® Complete

MAXIMUM STRENGTH

Menstrual Complete

Acetaminophen, Caffeine, Pyrilamine maleate

PAIN RELIEVER / DIURETIC / ANTIHISTAMINE

FOR MULTI-SYMPTOM RELIEF OF:
Cramps ι Bloating ι Fatigue
Backache ι Headache

16 Caplets

ACTUAL SIZE

TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING

*This product is not manufactured or distributed by Bayer HealthCare LLC, owner of the registered trademark Midol® Complete.

50844 REV0517A39021

Distributed by
LNK INTERNATIONAL, INC.
60 Arkay Drive, Hauppauge, NY 11788
USA

QUALITY PLUS

NDC 50844-390-21

*Compare to the active ingredients in Midol® Complete

MAXIMUM STRENGTH

Menstrual Complete

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No print /No varnish
Lot no/exp date

Drug Facts (continued)

■ TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN ■ see end flap for expiration date and lot number

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

B-1603-390-21-R
REV0517A39021

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.

Directions ■ do not take more than the recommended dose

■ adults and children 12 years and over:
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■ repeat every 6 hours, as needed
■ do not exceed 6 caplets per day
■ children under 12 years: ask a doctor

Other information ■ store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)

Drug Facts (continued)

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

Drug Facts (continued)

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

When using this product
■ alcohol, sedatives, and tranquilizers may increase drowsiness ■ you may get drowsy
■ excitability may occur, especially in children
■ be careful 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 heartbeat. The recommended dose of this product contains about as much caffeine as a cup of coffee. ■ avoid alcoholic beverages

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.

Drug Facts (continued)

symptoms associated with menstrual periods:
■ bloating ■ headache ■ muscle aches
■ cramps ■ backache ■ fatigue
■ water-weight gain

Warnings

Liver warning: This product contains acetaminophen. Severe liver damage may occur

Drug Facts

KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION

Active ingredients (in each caplet)	Purpose
Acetaminophen 500 mg	Pain reliever
Caffeine 60 mg	Diuretic
Pyrilamine maleate 15 mg	Antihistamine

Uses for the temporary relief of these

No print area
Glue flap

No print area
Glue flap

Quality Plus 44-390

MENSTRUAL COMPLETE MAXIMUM STRENGTH

acetaminophen, caffeine and pyrilamine maleate tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50844-390
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	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: M28OL1HH48)	
CROSPOVIDONE, UNSPECIFIED (UNII: 2S7830E561)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
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			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50844-390-19	1 in 1 CARTON	04/29/2002	
1		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:50844-390-	2 in 1 CARTON	04/29/2002	

2	21	2 in 1 CARTON	04/29/2002	
2		8 in 1 BLISTER PACK; 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	04/29/2002	

Labeler - L.N.K. International, Inc. (038154464)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(50844-390)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(50844-390) , pack(50844-390)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(50844-390)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(50844-390)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(50844-390)

Revised: 8/2023 L.N.K. International, Inc.