

RAPID RELEASE PAIN RELIEF- acetaminophen tablet, film coated
Publix Super Markets Inc

Publix Super Markets, Inc. Rapid Release Pain Relief Drug Facts

Active ingredient (in each caplet)

Acetaminophen 500 mg

Purpose

Pain reliever/fever reducer

Uses

- temporarily relieves minor aches and pains due to:
- headache
- muscular aches
- backache
- minor pain of arthritis
- the common cold
- toothache
- premenstrual and menstrual cramps
- temporarily reduces fever

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

liver disease

Ask a doctor or pharmacist before use if you are

taking the blood thinning drug warfarin

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.

Overdose warning: In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). Quick 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 (see overdose warning)

adults and children 12 years and over	<ul style="list-style-type: none">• take 2 caplets every 6 hours while symptoms last• do not take more than 6 caplets in 24 hours, unless directed by a doctor• do not use for more than 10 days unless directed by a doctor
children under 12 years	ask a doctor

Inactive ingredients

croscarmellose sodium, crospovidone, FD&C red #40 aluminum lake, FD&C yellow #6 aluminum lake, hypromellose, mica-based pearlescent pigment, polyethylene glycol, polysorbate 80, povidone, pregelatinized starch, stearic acid

Package/Label Principal Display Panel

EXTRA STRENGTH

rapid release pain relief

ACETAMINOPHEN 500 mg

Pain reliever/fever reducer

Fast relief

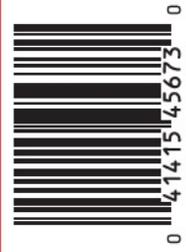
For adults

ACTUAL SIZE

50 CAPLETS

Compare to Tylenol® Extra Strength Rapid Release Gels active ingredient

3S071 63 C3



Drug Facts

Active ingredient (in each caplet)
Acetaminophen 500 mg...Pain reliever/fever reducer

Purpose

Uses ■ temporarily relieves minor aches and pains due to:

- the common cold
- minor pain of arthritis
- headache
- muscular aches
- backache
- toothache
- menstrual and menstrual cramps
- temporarily reduces fever

Warnings

Liver warnings: 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 redness ■ 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 liver disease

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

Drug Facts (continued)

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
- rashness 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. Overdose warning- In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). Quick 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 (see overdose warning)

adults and children 12 years and over

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- do not use for more than 10 days unless directed by a doctor

children under 12 years

- ask a doctor

Inactive ingredients croscarmellose sodium, croscopolone, FD&C red #40 aluminum lake, FD&C yellow #6 aluminum lake, hypromellose, meta-based pearlescent pigment, polyethylene glycol, polysochate 80, pov-done, pregelatinized starch, stearic acid

Publix
COMPLETE SATISFACTION OR YOUR MONEY BACK
PUBLIX GUARANTEE:
1-888-267-3037
LAKELAND, FL 33811 publix.com

DO NOT USE IF PRINTED SEAL UNDER CAP IS BROKEN OR MISSING
Store at 20-25°C (68-77°F).

NDC 56062-650-71



EXTRA STRENGTH
**rapidrelease
painrelief**

ACETAMINOPHEN 500 mg
Pain reliever/fever reducer

- Fast relief
- For adults



ACTUAL SIZE

50 CAPLETS

Compare to Tylenol® Extra Strength Rapid Release Gels active ingredient!

NDC 56062-650-71



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RAPID RELEASE PAIN RELIEF
acetaminophen tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:56062-650
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg

Inactive Ingredients

Ingredient Name	Strength
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
CROSPVIDONE (120 .MU.M) (UNII: 68401960MK)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

Product Characteristics

Color	RED	Score	no score
Shape	OVAL	Size	18mm
Flavor		Imprint Code	3S0
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:56062-650-71	1 in 1 CARTON	03/25/2016	
1		50 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:56062-650-78	1 in 1 CARTON	03/25/2016	
2		100 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 Drug	M013	03/25/2016	

Labeler - Publix Super Markets Inc (006922009)