

**ACETAMINOPHEN- acetaminophen tablet
LEADER/ Cardinal Health 110, Inc.**

5427-Cardinal

Drug Facts

Active ingredient

Acetaminophen 500 mg

Purpose

Pain Reliever/ Fever Reducer

Uses

Temporarily reduces fever and relieves minor aches and pains due to:

- headache
- muscular aches
- common cold
- toothache
- backache
- minor pain of arthritis
- premenstrual and menstrual cramps

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 are allergic to acetaminophen or any of the inactive ingredients in this product

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:

- take 2 tablets every 6 hours while symptoms last
- do not take more than 6 tablets in 24 hours, unless directed by a doctor
- do not take for more than 10 days unless directed by a doctor

children under 12 years: ask a doctor

Other Information

store at room temperature

Povidone, pregelatinized starch, sodium starch glycolate, stearic acid

Questions or comments?

Call (800) 231-4670

*This product is not manufactured or distributed by McNeil Consumer Healthcare, owner of the registered trademark Tylenol® Extra Strength.

DISTRIBUTED BY CARDINAL HEALTH, DUBLIN, OHIO 43017

www.myleader.com 1-800-200-6313

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

100% Money Back Guarantee

Return to place of purchase if not satisfied.

LEADER™

NDC 70000-0410-2

Extra Strength **Acetaminophen**

Tablets, 500 mg

Pain Reliever/ Fever Reducer

For Adults

Compare to Tylenol® Extra Strength active ingredient*

100 Tablets

8004118
(OF # & EXP. DATE)

NDC 70000-0410-2

LEADER²

**Extra Strength
Acetaminophen**

Tablets, 500 mg
Pain Reliever / Fever Reducer
FOR ADULTS

Actual Size
100 TABLETS

**COMPARE TO
TYLENOL®
EXTRA
STRENGTH
active ingredient***

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

Drug Facts	Purpose
Active ingredient (in each tablet) Acetaminophen 500 mg • Pain Reliever/ Fever Reducer	
Uses Temorarily reduces fever and relieves minor aches and pains due to: • headache • muscular aches • common cold • toothache • backache • minor pain of arthritis • premenstrual and menstrual cramps	
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 are allergic to acetaminophen or any of the inactive ingredients in this product ▲	

© 2018 Cardinal Health **Drug Facts** (continued under label)

CIN 5457 262

REV. 11/18

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PEEL HERE FOR MORE INFORMATION

Drug Facts (continued)

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: take 2 tablets every 6 hours while symptoms last
- do not take more than 6 tablets in 24 hours, unless directed by a doctor
- do not take for more than 10 days unless directed by a doctor
- children under 12 years: ask a doctor

Other Information

store at room temperature

Inactive Ingredients

Povidone, pregelatinized starch, sodium starch glycolate, stearic acid

Questions or comments?

Call (800) 231-4670

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100% Money Back Guarantee
Return to place of purchase if not satisfied.

NO COPY
ADHESIVE ZONE

EZ-OPEN CAP

LEADER²

NDC70000-0036-1

Extra Strength
Acetaminophen

Tablets, 500 mg
Pain Reliever / Fever Reducer

FOR ADULTS
100 TABLETS

COMPARE TO
TYLENOL[®]
EXTRA STRENGTH
active ingredient*

THIS PACKAGE FOR HOUSEHOLDS WITHOUT YOUNG CHILDREN

15406-05-19
LOT # & EXP. DATE

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

Drug Facts

Active ingredient (in each tablet)
Acetaminophen 500 mg. Pain Reliever / Fever Reducer

Uses

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

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 are allergic to acetaminophen or any of the inactive ingredients in this product

© 2019 Cardinal Health Drug Facts (continued under label)

CIN 5528138 REV. 5/19



PEEL HERE FOR MORE INFORMATION

Drug Facts (continued)

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: take 2 tablets every 6 hours while symptoms last
- do not take more than 6 tablets in 24 hours, unless directed by a doctor
- do not take for more than 10 days unless directed by a doctor

children under 12 years: ask a doctor

Other information

store at room temperature

Inactive ingredients

Povidone, pregelatinized starch, sodium starch glycolate, stearic acid

Questions or comments?

Call (800) 231-4670

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www.tylenol.com 1-800-200-6337

100% Money Back Guarantee

Return to place of purchase if not satisfied.

LEADER²

NDC70000-0094

Extra Strength
Acetaminophen

Tablets, 500 mg
Pain Reliever / Fever Reducer

For Adults

500 TABLETS

Actual Size

COMPARE TO
EXTRA STRENGTH
TYLENOL[®]
active ingredient*

100% Money Back Guarantee

Drug Facts

Active ingredient (in each tablet) Purpose
Acetaminophen 500 mg Pain reliever/fever reducer

Uses ■ temporarily relieves minor aches and pains due to ■ the common cold ■ headache ■ toothache ■ muscular aches ■ backache ■ minor pain of arthritis ■ 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 are allergic to acetaminophen or any of the inactive ingredients in this product

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.

Drug Facts (continued)

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 of age and over
 - take 2 tablets every 6 hours while symptoms last
 - do not take more than 6 tablets in 24 hours, unless directed by a doctor
 - do not take for more than 10 days unless directed by a doctor
- children under 12 years of age
 - ask a doctor

Other information ■ Store in a dry place at 15° - 30°C (59° - 86°F)

Inactive ingredients povidone, pregelatinized starch, sodium starch glycolate and stearic acid

Questions or comments? Call 1-800-231-4670

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING. R53427

*This product is not manufactured or distributed by Kenvue Inc., owner of the registered trademark Tylenol®

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www.tylenol.com 1-800-200-6337

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CIN 5933641 REV. 07/24



ACETAMINOPHEN
acetaminophen tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70000-0709
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
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
STARCH, CORN (UNII: O8232NY3SJ)	
POVIDONE K30 (UNII: U725QWY32X)	

Product Characteristics

Color	white	Score	2 pieces
Shape	ROUND	Size	12mm
Flavor		Imprint Code	54;27
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70000-0709-1	500 in 1 BOTTLE; Type 0: Not a Combination Product	12/10/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	12/10/2024	

ACETAMINOPHEN

acetaminophen tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70000-0410
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		
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)				
STEARIC ACID (UNII: 4ELV7Z65AP)				
STARCH, CORN (UNII: O8232NY3SJ)				
POVIDONE K30 (UNII: U725QWY32X)				
Product Characteristics				
Color	white	Score	2 pieces	
Shape	ROUND	Size	12mm	
Flavor		Imprint Code	54;27	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70000-0410-2	100 in 1 BOTTLE; Type 0: Not a Combination Product	11/05/2018	
2	NDC:70000-0410-1	1 in 1 CARTON	12/19/2018	
2		60 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	11/05/2018		

ACETAMINOPHEN			
acetaminophen tablet			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70000-0036
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
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
STARCH, CORN (UNII: O8232NY3SJ)	
POVIDONE K30 (UNII: U725QWY32X)	

Product Characteristics

Color	white	Score	2 pieces
Shape	ROUND	Size	12mm
Flavor		Imprint Code	54;27
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70000-0036-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	05/28/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	05/28/2019	

Labeler - LEADER/ Cardinal Health 110, Inc. (063997360)

Revised: 1/2025

LEADER/ Cardinal Health 110, Inc.