

ACETAMINOPHEN- acetaminophen tablet
Bryant Ranch Prepack

5427-Major

Drug Facts

Active Ingredient (in each tablet)

Acetaminophen 500 mg

Purpose

Pain reliever/ Fever reducer

Uses

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

- headache
- muscular aches
- backache
- minor pain of arthritis
- the common cold
- toothache
- 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.

- 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 in a dry place at 15° – 30°C (59° – 86°F).

Povidone, pregelatinized starch, sodium starch glycolate, stearic acid

Questions or comments?

Call 1-800-231-4670

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

Distributed by:

**MAJOR® PHARMACEUTICALS
Indianapolis, IN 46268**

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

HOW SUPPLIED

NDC: 72162-2164-1: 100 Tablets in a BOTTLE

Repackaged/Relabeled by:
Bryant Ranch Prepack, Inc.
Burbank, CA 91504

Acetaminophen 500mg Tablets #100



GTIN 00363629106424
 Lot LOT123
 Exp 06/02/2025
 SN 1234567890

Drug Facts	
Active ingredient (in each tablet)	Purpose
Acetaminophen 500 mg	Pain reliever/Fever reducer
Uses	
Temporarily reduces fever and relieves minor aches and pains due to: •headache •muscular aches •backache •minor pain of arthritis •common cold •toothache •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.	
•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.	
Other Information	
•Store at room temperature. •Do not use if imprinted safety seal under cap is broken or missing. •Aspirin Free •Extra Strength •Pain Reliever	
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	
Inactive Ingredients	
Povidone, pregelatinized starch, sodium starch glycolate, stearic acid.	

NDC 72162-2164-1

Acetaminophen Tablets

500 mg



Relabeled by:
 Bryant Ranch Prepack, Inc.
 Burbank, CA 91504 USA

100 Tablets
 Manufactured by:
 Major Pharmaceuticals



7216221641

ACETAMINOPHEN

acetaminophen tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72162-2164(NDC:0904-6730)
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
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
POVIDONE K30 (UNII: U725QWY32X)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start	Marketing End
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#	Item Code	Package Description	Date	Date
1	NDC:72162-2164-1	1 in 1 CARTON	09/12/2018	
1		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	09/12/2018	

Labeler - Bryant Ranch Prepack (171714327)

Registrant - Bryant Ranch Prepack (171714327)

Establishment

Name	Address	ID/FEI	Business Operations
Bryant Ranch Prepack		171714327	REPACK(72162-2164) , RELABEL(72162-2164)

Revised: 5/2024

Bryant Ranch Prepack