

ACETAMINOPHEN- acetaminophen tablet

Bryant Ranch Prepack

Major Pharmaceuticals Acetaminophen Drug Facts

Active ingredient (in each caplet)

Acetaminophen 500 mg

Purpose

Pain reliever/fever reducer

Uses

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

carnauba wax, corn starch*, croscarmellose sodium*, hypromellose, polyethylene glycol, povidone, pregelatinized starch, sodium starch glycolate*, stearic acid

*may contain one or more of these ingredients

Questions or comments?

1-800-616-2471

HOW SUPPLIED

NDC: 71335-2251-1: 30 TABLETs in a BOTTLE

Acetaminophen 500 mg Caplet



GTIN 00371335225117
 Lot 208620
 Exp 1/29/2026
 SN 0123456789

Drug Facts	
Active ingredient (in each caplet)	Purpose
Acetaminophen 500 mg	Pain reliever/fever reducer
Uses	
•temporarily relieves minor aches and pains due to -the common cold -headache -backache -minor pain of arthritis -toothache -muscular aches -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 redness -hives -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.	
Other Information	
•Store at 20-25°C (68-77°F).	
•Do not use if printed seal under cap is broken or missing.	
•ASPIRIN FREE	
•DONOT USE WITH OTHER MEDICINES CONTAINING ACETAMINOPHEN	
•PAIN RELIEVER/REDUCER	
•FOR ADULTS	
Directions	
•do not take more than directed (see overdose warning). adults and children 12 years and over -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	
carnauba wax, corn starch*, croscarmellose sodium*, hypromellose, polyethylene glycol, povidone, pregelatinized starch, sodium starch glycolate*, stearic acid	
*may contain one or more of these ingredients	

NDC 71335-2251-1

Acetaminophen Caplets

500 mg

30 Caplets



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

Manufactured by:
 Major
 Pharmaceuticals



ACETAMINOPHEN			
acetaminophen tablet			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71335-2251(NDC:0904-6720)
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
CARNAUBA WAX (UNII: R12CBM0EIZ)			
STARCH, CORN (UNII: O8232NY3S)			
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)			
STEARIC ACID (UNII: 4ELV7Z65AP)			
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)			

Product Characteristics

Color	WHITE	Score	no score
Shape	OVAL	Size	16mm
Flavor		Imprint Code	L484
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71335-2251-1	30 in 1 BOTTLE; Type 0: Not a Combination Product	01/29/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	07/26/2018	

Labeler - Bryant Ranch Prepack (171714327)

Registrant - Bryant Ranch Prepack (171714327)

Establishment

Name	Address	ID/FEI	Business Operations
Bryant Ranch Prepack		171714327	REPACK(71335-2251) , RELABEL(71335-2251)

Revised: 1/2024

Bryant Ranch Prepack