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

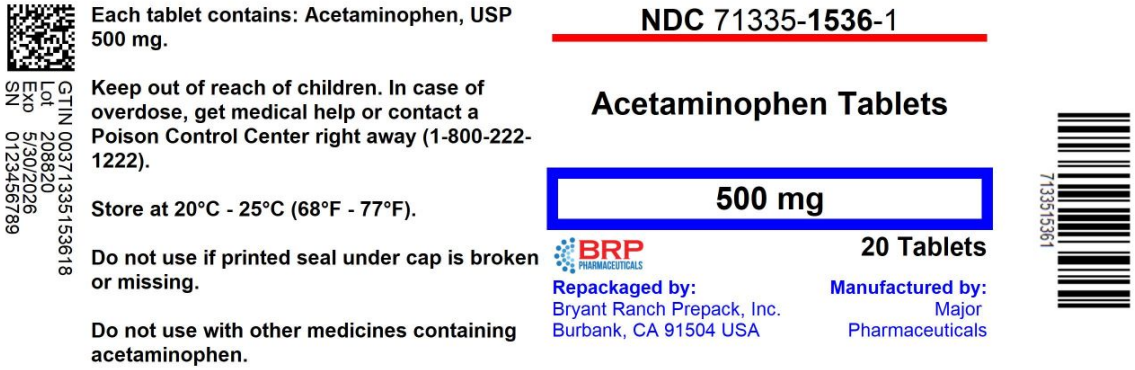
Acetaminophen Tablets 500 mg

- NDC: 71335-1536-1: 20 Tablets in a BOTTLE
- NDC: 71335-1536-2: 15 Tablets in a BOTTLE
- NDC: 71335-1536-3: 40 Tablets in a BOTTLE
- NDC: 71335-1536-4: 100 Tablets in a BOTTLE

- NDC: 71335-1536-5: 30 Tablets in a BOTTLE
- NDC: 71335-1536-6: 45 Tablets in a BOTTLE
- NDC: 71335-1536-7: 50 Tablets in a BOTTLE
- NDC: 71335-1536-8: 60 Tablets in a BOTTLE
- NDC: 71335-1536-9: 90 Tablets in a BOTTLE
- NDC: 71335-1536-0: 250 Tablets in a BOTTLE

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

Acetaminophen Tablets 500 mg



ACETAMINOPHEN

acetaminophen tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71335-1536(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 Date	Marketing End Date
1	NDC:71335-1536-1	20 in 1 BOTTLE; Type 0: Not a Combination Product	05/30/2024	
2	NDC:71335-1536-2	15 in 1 BOTTLE; Type 0: Not a Combination Product	05/30/2024	
3	NDC:71335-1536-3	40 in 1 BOTTLE; Type 0: Not a Combination Product	05/30/2024	
4	NDC:71335-1536-4	100 in 1 BOTTLE; Type 0: Not a Combination Product	04/17/2020	
5	NDC:71335-1536-5	30 in 1 BOTTLE; Type 0: Not a Combination Product	06/16/2020	
6	NDC:71335-1536-6	45 in 1 BOTTLE; Type 0: Not a Combination Product	05/30/2024	
7	NDC:71335-1536-7	50 in 1 BOTTLE; Type 0: Not a Combination Product	05/30/2024	
8	NDC:71335-1536-8	60 in 1 BOTTLE; Type 0: Not a Combination Product	05/30/2024	
9	NDC:71335-1536-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	05/30/2024	
10	NDC:71335-1536-0	250 in 1 BOTTLE; Type 0: Not a Combination Product	05/30/2024	

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(71335-1536) , RELABEL(71335-1536)