EXTRA STRENGTH PAIN RELIEF- acetaminophen tablet
Bryant Ranch Prepack

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gc201

Active Ingredients

Acetaminophen 500 mg

Purpose

Pain Reliever/Fever Reducer

Uses

- temporarily relieves minor aches and pains
- temporarily reduces fever

Warnings

Liver warning:

This product contains acetaminophen. Severe liver damage may occur if you take

- more than 8 tablets (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.

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 symptom 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. In case of overdose, get medical help or contact a Poison Control Center
right away. 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**
- adults and children 12 years and over: Take 1 to 2 tablets every 4 to 6 hours, as needed; not more than 6 tablets in 24 hours. Do not take for more than 10 days unless directed by a doctor
- children under 12 years: ask a doctor

Other Information

- **TAMPER EVIDENT: Do not use if imprinted seal under cap is missing or broken.**
- store at 20ºC-25ºC (68ºF-77ºF)

Inactive Ingredients

povidone, sodium starch glycolate, starch, stearic acid.

Questions or comments?

1-800-540-3765

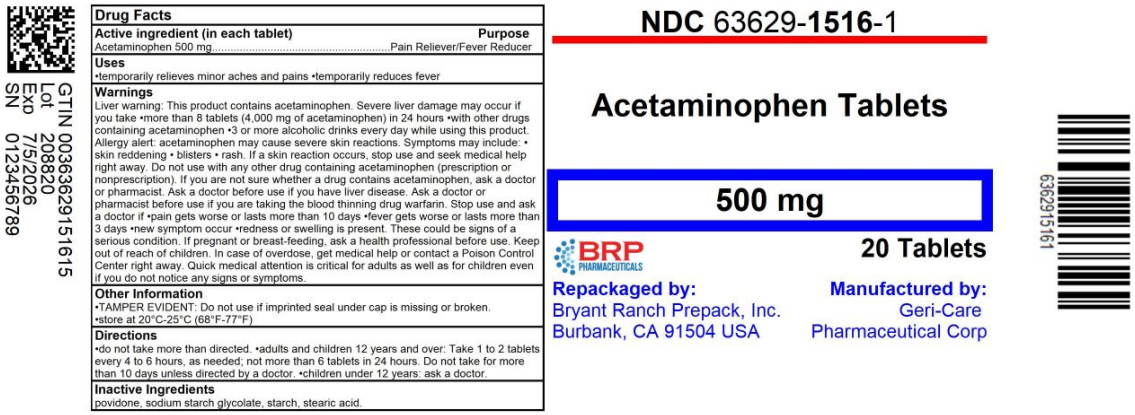
HOW SUPPLIED

Acetaminophen 500 mg Tablet

- NDC: 63629-1516-1: 20 Tablets in a BOTTLE
- NDC: 63629-1516-2: 15 Tablets in a BOTTLE
- NDC: 63629-1516-3: 40 Tablets in a BOTTLE
- NDC: 63629-1516-4: 100 Tablets in a BOTTLE
- NDC: 63629-1516-5: 30 Tablets in a BOTTLE
- NDC: 63629-1516-6: 45 Tablets in a BOTTLE
- NDC: 63629-1516-7: 50 Tablets in a BOTTLE
- NDC: 63629-1516-8: 60 Tablets in a BOTTLE
- NDC: 63629-1516-9: 90 Tablets in a BOTTLE
- NDC: 63629-1516-0: 250 Tablets in a BOTTLE

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

Acetaminophen 500 mg Tablet



EXTRA STRENGTH PAIN RELIEF

acetaminophen tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63629-1516(NDC:57896-201)
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)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics			
Color	white (WHITE)	**Score**	no score
Shape	ROUND (Round)	**Size**	12mm
Flavor		**Imprint Code**	M2A457344
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**

1	NDC:63629-1516-1	20 in 1 BOTTLE; Type 0: Not a Combination Product	06/17/2008	
2	NDC:63629-1516-2	15 in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2023	
3	NDC:63629-1516-3	40 in 1 BOTTLE; Type 0: Not a Combination Product	12/29/2004	
4	NDC:63629-1516-4	100 in 1 BOTTLE; Type 0: Not a Combination Product	02/21/2006	
5	NDC:63629-1516-5	30 in 1 BOTTLE; Type 0: Not a Combination Product	03/27/2007	
6	NDC:63629-1516-6	45 in 1 BOTTLE; Type 0: Not a Combination Product	08/11/2009	
7	NDC:63629-1516-7	50 in 1 BOTTLE; Type 0: Not a Combination Product	09/10/2008	
8	NDC:63629-1516-8	60 in 1 BOTTLE; Type 0: Not a Combination Product	10/09/2008	
9	NDC:63629-1516-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	12/21/2010	
10	NDC:63629-1516-0	250 in 1 BOTTLE; Type 0: Not a Combination Product	09/04/2009	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	01/01/1989	

Labeler - Bryant Ranch Prepack (171714327)

Registrant - Bryant Ranch Prepack (171714327)

Establishment

Name	Address	ID/FEI	Business Operations
Bryant Ranch Prepack		171714327	REPACK(63629-1516) , RELABEL(63629-1516)

Revised: 7/2024 Bryant Ranch Prepack