

**ACETAMINOPHEN- acetaminophen tablet
Bryant Ranch Prepack**

**Acetaminophen 325 mg Tablets
Regular Strength**

Active ingredient (in each tablet)

Acetaminophen 325 mg

Purpose

Pain Reliever/Fever Reducer

Uses

To reduce fever and for the temporary relief of minor aches and pains due to:

- headache
- muscular aches
- backache
- minor pain of arthritis
- the common cold
- toothache
- premenstrual and menstrual cramps.
- Temporarily reduces fever.

Warnings

Liver warning:

This product contains acetaminophen. Severe liver damage may occur if

- adult takes more than 4,000 mg of acetaminophen in 24 hours
- child takes more than 5 doses in 24 hours
- taken with other drugs containing acetaminophen
- adult has 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 non-prescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- if the user has ever had an allergic reaction to this product or any of its ingredients.

Ask a doctor before use if the user has

- has liver disease
- is a child with pain of arthritis

Ask a doctor or pharmacist before use if your child is taking the blood thinning drug warfarin

Stop use and ask a doctor if:

- pain gets worse or lasts more than 10 days (for adults) or 5 days (for children)
- 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:

Taking more than the recommended dose (overdose) may cause liver damage. 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 & children 12 years and over	<ul style="list-style-type: none">• take 2 tablets every 4 to 6 hours while symptoms last• do not take more than 10 tablets in 24 hours• do not use for more than 10 days unless directed by a doctor
children 6-11 years	<ul style="list-style-type: none">• take 1 tablet every 4 to 6 hours while symptoms last• do not take more than 5 tablets in 24 hours• do not use for more than 5 days unless directed by a doctor
child under 6 years	ask a doctor

Other information

- store at 20° to 25°C (68° to 77°F)

Inactive Ingredients:

Gelatinized starch, magnesium stearate, povidone

Questions or Comments?

Call toll-free 1-877-225-6999

**Manufactured for
Akron Pharma, Inc.,**

373 RT US46 W Building E,

Suite 117, Fairfield, NJ - 07004

Mfg. Lic. No: TN/DRUGS/558/1997

* This product is not manufactured or distributed by Johnson and Johnson, consumer inc., distributor of regular Tylenol Tablets.

HOW SUPPLIED

Acetaminophen 325 mg tablet: white color, round shape and the imprint code is A325.

NDC: 71335-2747-1: 20 Tablets in a BOTTLE

NDC: 71335-2747-2: 100 Tablets in a BOTTLE

NDC: 71335-2747-3: 30 Tablets in a BOTTLE

NDC: 71335-2747-4: 2 Tablets in a BOTTLE

NDC: 71335-2747-5: 6 Tablets in a BOTTLE

NDC: 71335-2747-6: 10 Tablets in a BOTTLE

NDC: 71335-2747-7: 60 Tablets in a BOTTLE

NDC: 71335-2747-8: 90 Tablets in a BOTTLE

NDC: 71335-2747-9: 40 Tablets in a BOTTLE

NDC: 71335-2747-0: 24 Tablets in a BOTTLE

Store at 20° to 25°C (68° to 77°F)

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

Acetaminophen 325 mg Tablet



GTIN 0037135274719
 Lot 208820
 Exp 10/2/2027
 SN 0123456789

Drug Facts	
Active ingredient (in each tablet) Acetaminophen 325 mg	Purposes Pain Reliever/Fever Reducer
Uses temporarily relieves minor aches and pains temporarily reduces fever	
Warnings 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 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.	
Other Information Tamper Evident: Do not use if imprinted seal under cap is missing or broken. Store at 20°C-25°C (68°F-77°F). Scan Package Insert QR Code for more information.	
Directions do not take more than directed. adults and children 12 years and over: Take 2 tablets every 4-6 hours, as needed; not more than 10 tablets in 24 hours. Do not take for more than 10 days unless directed by a doctor. children under 12 years: ask a doctor.	
Inactive Ingredients povidone, sodium starch glycolate, starch, stearic acid.	

NDC 71335-2747-1

Acetaminophen Tablets

325 mg

20 Tablets



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

Manufactured by:
Akron Pharma Inc.



Package Insert

ACETAMINOPHEN

acetaminophen tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71335-2747(NDC:71399-1025)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg

Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	

Product Characteristics

Color	white	Score	2 pieces
Shape	ROUND	Size	4mm
Flavor		Imprint Code	A325
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71335-2747-1	20 in 1 BOTTLE; Type 0: Not a Combination Product	10/02/2025	
2	NDC:71335-2747-2	100 in 1 BOTTLE; Type 0: Not a Combination Product	10/02/2025	
3	NDC:71335-2747-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	10/02/2025	

4	NDC:71335-2747-4	2 in 1 BOTTLE; Type 0: Not a Combination Product	10/02/2025	
5	NDC:71335-2747-5	6 in 1 BOTTLE; Type 0: Not a Combination Product	10/02/2025	
6	NDC:71335-2747-6	10 in 1 BOTTLE; Type 0: Not a Combination Product	10/02/2025	
7	NDC:71335-2747-7	60 in 1 BOTTLE; Type 0: Not a Combination Product	10/02/2025	
8	NDC:71335-2747-8	90 in 1 BOTTLE; Type 0: Not a Combination Product	10/02/2025	
9	NDC:71335-2747-9	40 in 1 BOTTLE; Type 0: Not a Combination Product	10/02/2025	
10	NDC:71335-2747-0	24 in 1 BOTTLE; Type 0: Not a Combination Product	10/02/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	12/06/2024	

Labeler - Bryant Ranch Prepack (171714327)

Registrant - Bryant Ranch Prepack (171714327)

Establishment

Name	Address	ID/FEI	Business Operations
Bryant Ranch Prepack		171714327	REPACK(71335-2747) , RELABEL(71335-2747)

Revised: 10/2025

Bryant Ranch Prepack