#### ACETAMINOPHEN- acetaminophen tablet Walmart

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Pain Reliever Acetaminophen USP, 500 mg Pain Reliever/Fever Reducer Contains No Aspirin

# Active ingredient (in each gelcap)

Acetaminophen USP, 500 mg

#### Purpose

Pain reliever/fever reducer

#### Uses

- temporarily relieves minor aches and pains due to:
- headache
- muscular aches
- backache
- minor pain of arthritis
- the common cold n toothache
- premenstrual and menstrual cramps
- temporarily reduces fever

### Liver warning:

This product contains acetaminophen. Severe liverdamage may occur if you take

- more than 4,000mg 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 seekmedical 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 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 gelcaps every 6 hours while symptoms last
- do not take more than 6 gelcaps 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

### Other information

- store at 20°-25°C (68°-77°F)
- avoid high humidity
- see bottom of the label for expiration date and lot number

### Inactive ingredients

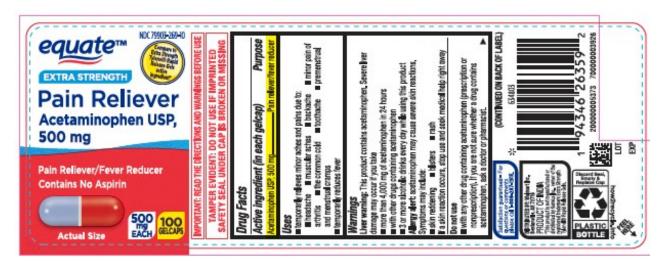
ammonium hydroxide, black iron oxide, colloidal silicon dioxide, croscarmellose sodium, D&C red #33, FD&C blue #1, FD&C red #40,

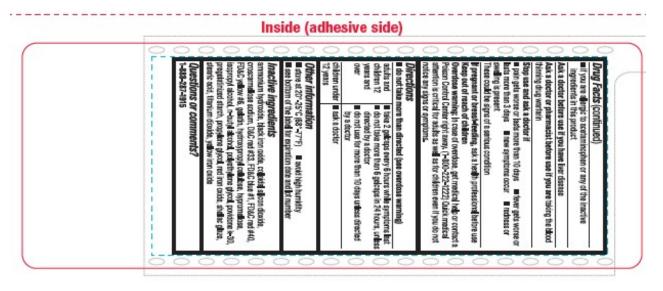
FD&C yellow #6, gelatin, hydroxypropyl cellulose, hypromellose, isopropyl alcohol, nbutyl alcohol, polyethylene glycol, povidone k-30,

pregelatinized starch, propylene glycol, red iron oxide, shellac glaze, stearic acid, titanium dioxide, yellow iron oxide

### Questions or comments?

1-888-287-1915







Inside (adhesive side)
Trug Facts (Continued)   Regeneration of existence   Control Contents and three tend (see overdose warning)   activation of existence   Content tend metered (see overdose warning)   activation   Exists a concer   activation is concer   activation   Exists a concer   active and one state of the exists man to days unless directed by a doctor   The information   active information

ACETAMINOPHEN					
acetaminophen tablet					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Sou	ırce)	NDC:799	03-269
Route of Administration	ORAL				
A					
Active Ingredient/Active	моюту				
Ingredient Name Basis of Streng				rength	Strength
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) ACETAMINOPH			EN	500 mg	
Inactive Ingredients					
mactive myrealents					
Ingredient Name			S	Strength	
AMMONIA (UNII: 5138Q19F1X)					
HYDROXYPROPYL CELLULOSE, U	JNSPECIFIED (UNII: 9XZ8H	16N6OH)			
CROSCARMELLOSE SODIUM (UN	II: M28OL1HH48)				

D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
GELATIN (UNII: 2G86QN327L)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
BUTYL ALCOHOL (UNII: 8PJ61P6TS3)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
STARCH, CORN (UNII: 08232NY3SJ)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POVIDONE K30 (UNII: U725QWY32X)	
FD&C RED NO. 40 (UNII: WZ B9127XOA)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
FERRIC OXIDE YELLOW (UNII: EX43802MRT)	
SHELLAC (UNII: 46N107B710)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Product Characteristics				
Color	red (red, blue (blue-gray))	Score	no score	
Shape	CAPSULE (banded two-part capsule)	Size	19mm	
Flavor		Imprint Code	G;1	
Contains				

# Packaging

1   NDC:79903-269- 10   100 in 1 BOTTLE; Type 0: Not a Combination Product   01/16/2025     2   NDC:79903-269- 22   225 in 1 BOTTLE; Type 0: Not a Combination Product   01/16/2025	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1			01/16/2025	
	2			01/16/2025	

# Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
OTC Monograph Drug	M013	01/16/2025	

# Labeler - Walmart (051957769)

Revised: 1/2025

Walmart