

ACETAMINOPHEN- acetaminophen tablet
Walmart

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, n-butyl 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

equate™ NDC 79903-269-40
EXTRA STRENGTH
Pain Reliever
Acetaminophen USP,
500 mg
 Pain Reliever/Fever Reducer
 Contains No Aspirin
 Actual Size **500 mg EACH** **100 GELCAPS**

Compare It To
 To know how
 much better
 Equate is

IMPORTANT: READ THE DIRECTIONS AND WARNINGS BEFORE USE
TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

Drug Facts
Active ingredient (in each gelcap) Purpose
 Acetaminophen USP, 500 mg Pain reliever/fever reducer

Uses
 Temporarily relieves 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 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
 ■ 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.

(CONTINUED ON BACK OF LABEL)
 634603
 1 94346 26359 2
 200000005373 7000000005956
 LOT EXP
 Satisfaction guaranteed™ For questions or concerns call 1-888-287-1915
 634603
 Manufactured by:
 Watson Laboratories, Inc.
 200000005373
PRODUCT OF INDIA
 This product is manufactured by Watson Laboratories, Inc. through its wholly owned subsidiary, Watson Pharmaceuticals, Inc. All rights reserved.
 PLASTIC BOTTLE
 100% RECYCLED
 100% RECYCLED

Inside (adhesive side)

Drug Facts (continued)
 ■ If you are allergic to acetaminophen or any of the inactive ingredients in this product.
 Ask a doctor or pharmacist before use if you have liver disease, kidney or lung problems.
 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, 1-800-272-1222. 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 with symptoms that 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 bottle 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, n-butyl 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

D&C RED NO. 33 (UNII: 9DBA05BB0L)
FERROSFERRIC 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: O8232NY3SJ)
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
POVIDONE K30 (UNII: U725QWY32X)
FD&C RED NO. 40 (UNII: WZB9127XOA)
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)
ISOPROPYL ALCOHOL (UNII: ND2M416302)
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)
SHELLAC (UNII: 46N107B71O)
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

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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	

Marketing Information

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

Labeler - Walmart (051957769)