ACETAMINOPHEN- acetaminophen tablet, film coated, extended release WAL-MART STORES INC

Arthritis
Pain Reliever
ACETAMINOPHEN EXTENDED-RELEASE TABLETS USP, 650 mg
PAIN RELIEVER / FEVER REDUCER

ARTHRITIS 8 HOURCAPLETS

- For the temporary relief of minor arthritis pain
- Lasts up to 8 hours

Active ingredient

(in each caplet) Acetaminophen USP, 650 mg

Purpose

Pain reliever/fever reducer

Uses

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

Liver warning

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

- more than 6 caplets in 24 hours, which is the maximum daily amount
- 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
- 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

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

■ d o not take more than directed (see overdose warning)

adults

- take 2 caplets every 8 hours with water
- swallow whole; do not crush, chew, split or dissolve
- do not take more than 6 caplets in 24 hours
- do not use for more than 10 days unless directed by a doctor under 18 years of age
- ask a doctor

Other information

- store between 20-25°C (68-77°F)
- do not use if foil inner seal is broken or missing

Inactive ingredients

hydroxy ethyl cellulose, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, povidone, pregelatinized starch, sodium starch glycolate, stearic acid.

Questions or comments?

Contact 1-877-770-3183 Mon-Fri 8:00 AM EST to 5:00 PM PST.

PDP



ACETAMINOPHEN

acetaminophen tablet, film coated, extended release

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:49035-181

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII: 36209ITL9D) ACETAMINOPHEN 650 mg

Ingredient Name Strength HYDROXYETHYL CELLULOSE (140 MPA.S AT 5%) (UNII: 8136Y38GY5) HYPROMELLOSE 2910 (6 MPA.S) (UNII: 0WZ 8WG20P6) MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U) STARCH, CORN (UNII: 08232NY3SJ) POVIDONE K30 (UNII: U725QWY32X) SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)

STEARIC ACID (UNII: 4ELV7Z65AP)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	

Product Characteristics				
Color	white (White to off white colored)	Score	no score	
Shape	OVAL (Capsule shaped, biconvex intact film coated tablets)	Size	19mm	
Flavor		Imprint Code	G;650	
Contains				

l	P	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1	NDC:49035-181- 21	225 in 1 BOTTLE; Type 0: Not a Combination Product	10/04/2019		

Marketing I	arketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA211544	10/04/2019		

Labeler - WAL-MART STORES INC (051957769)

Revised: 12/2023 WAL-MART STORES INC