ACETAMINOPHEN - acetaminophen tablet, extended release WALMART INC.

Drug Facts

Active ingredient (in each extended-release tablet)

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 cramp toothache
 - the common cold
 - headache
 - toothache
- temporarily reduces fever

Warnings

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

- more than 6 tablets 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.

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:

- take 2 tablets every 8 hours with water
- swallow whole; do not crush, chew, split or dissolve
- do not take more than 6 tablets 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 at 20° to 25°C (68° to 77°F). Avoid excessive heat 40°C (104°F).
- USP Dissolution test is pending

Inactive ingredients

colloidal silicon dioxide, hydroxyethyl cellulose, hypromellose, magnesium stearate, microcrystalline cellulose, povidone, pregelatinized starch (maize), sodium starch glycolate, titanium dioxide, triacetin

Questions or comments? call 1-888-287-1915

DISTRIBUTED BY: Walmart Inc., Bentonville, AR 72716

Made in India Code: TS/DRUGS/22/2009

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 650 mg (325 Tablet Bottle)

TO OPEN: 1. PUSH DOWN 2. TURN CAP equate[™] NDC 79903-132-25

Compare to Tylenol[®] 8 HR Arthritis Pain active ingredient**

8 HOUR Arthritis Pain Relief Acetaminophen Extended-Release Tablets USP, 650 mg

Pain Reliever/Fever Reducer For the Temporary Relief of Minor Arthritis Pain

DO NOT USE WITH OTHER MEDICINES CONTAINING ACETAMINOPHEN

650	325
mg	Extended-Release
EACH	Tablets

Actual Size



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ACETAMINOPHEN

acetaminophen tablet, extended release

Product Information				
HUMAN OTC DRUG	Item Code (Sou	urce)	NDC:799	03-132
F Administration ORAL				
Active Ingredient/Active Moiety				
edient Name		Basis of St	rength	Strength
L9D) (ACETAMINOPHEN - UNI	I:362O9ITL9D)	ACETAMINOPH	EN	650 mg
Inactive Ingredients				
Ingredient Name			9	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				
40 MPA.S AT 5%) (UNII: 81	36Y38GY5)			
	ORAL Moiety redient Name L9D) (ACETAMINOPHEN - UNI Ingredient Name BU4)	ORAL Moiety redient Name L9D) (ACETAMINOPHEN - UNII: 36209ITL9D) Ingredient Name	ORAL Moiety redient Name Basis of St L9D) (ACETAMINOPHEN - UNII: 36209ITL9D) ACETAMINOPH Ingredient Name Busis BU4) Busis	ORAL Moiety redient Name Basis of Strength L9D) (ACETAMINOPHEN - UNII: 36209ITL9D) ACETAMINOPHEN

985)	
Score	no score
Size	19mm
Imprint Code	l;06
Marketing Start Date	Marketing End Date
07/26/2022	
h Marketing Start	Marketing End
Date	Date
	Date
	Size Imprint Code Marketing Start Date

Labeler - WALMART INC. (051957769)

Registrant - Aurohealth LLC (078728447)

Establishment			
Name	Address	ID/FEI	Business Operations
Aurobindo Pharma Limited		650381903	ANALYSIS(79903-132), MANUFACTURE(79903-132)

Revised: 1/2024

WALMART INC.