

ACETAMINOPHEN - acetaminophen tablet, extended release
KROGER COMPANY

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 cramps
 - 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).
- **do not use if carton is opened or foil inner seal is broken**
- Meets USP dissolution test 3

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?
1-800-632-6900

DISTRIBUTED BY THE KROGER CO.
CINCINNATI, OHIO 45202
MADE IN INDIA

Code: TS/DRUGS/22/2009

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 650 mg (24 Tablet Container Label)

NDC 30142-157-07

Kroger®

TO OPEN PUSH DOWN, TURN CAP

8 Hour

Arthritis Pain

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**

24

EXTENDED-

RELEASE TABLETS

Kroger NDC 30142-157-07
TO OPEN PUSH DOWN, TURN CAP
8 Hour
Arthritis Pain
Acetaminophen
Extended-Release
Tablets USP, 650 mg
Pain Reliever/Fever Reducer
For the Temporary Relief
of Minor Arthritis Pain
24
DO NOT USE WITH OTHER
MEDICINES CONTAINING
ACETAMINOPHEN
EXTENDED-
RELEASE TABLETS

DO NOT USE IF FOIL LINER SEAL IS BROKEN, CONTAINS NO ASPHIN

Active ingredient
(in each extended-release tablet)
Acetaminophen USP 650 mg — Pain reliever/fever reducer

Purpose
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

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. ■ You are urged to read the complete text of the package insert that comes with 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°). Avoid excessive heat.
40°C (104°F). ■ Meets USP dissolution test 3

Inactive ingredients: colloidal silicon dioxide, hydroxyethyl cellulose, hypromellose, magnesium stearate, microcrystalline cellulose, polydioxane, pregelatinized starch (maize), sodium starch glycolate, titanium dioxide, triacetin

Questions or comments? 1-800-632-6900
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Code: TS/DRUGS/22/2009

P 1432668 LM-5209
* Unvarnish zone

* Lot: XXXXXXXX
EXP: MM/YYYY
Prefix & Variables of Lot, EXP shall be printed online during packing.

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 650 mg (24 Tablets Container Carton)

**COMPARE TO the active ingredient of
TYLENOL® 8HR ARTHRITIS PAIN *See top panel**

NDC 30142-157-07

Kroger®

8 Hour

TO OPEN

PUSH DOWN,

TURN CAP

Arthritis Pain

**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**

actual size

**24 EXTENDED-
RELEASE TABLETS**



PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 650 mg (225 Tablet Container Label)

**COMPARE TO the active ingredient of
 TYLENOL® 8HR ARTHRITIS PAIN *See inside panel**

NDC 30142-157-35

EASY OPEN CAP

Kroger® 8 Hour

Arthritis Pain

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	650 mg

Inactive Ingredients

Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
HYDROXYETHYL CELLULOSE (140 MPA.S AT 5%) (UNII: 8136Y38GY5)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
STARCH, CORN (UNII: O8232NY3SJ)	
SODIUM STARCH GLYCOLATE TYPE B POTATO (UNII: 27NA468985)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	

Product Characteristics

Color	WHITE (White to Off-White)	Score	no score
Shape	CAPSULE (Caplet)	Size	19mm
Flavor		Imprint Code	I;06
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:30142-157-07	1 in 1 CARTON	05/25/2021	
1		24 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:30142-157-35	225 in 1 BOTTLE; Type 0: Not a Combination Product	05/25/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA207229	05/25/2021	

Labeler - KROGER COMPANY (006999528)**Registrant** - Aurohealth LLC (078728447)

Establishment

Name	Address	ID/FEI	Business Operations
Aurobindo Pharma Limited		650381903	ANALYSIS(30142-157) , MANUFACTURE(30142-157)

Revised: 1/2024

KROGER COMPANY