ACETAMINOPHEN- acetaminophen tablet, film coated, extended release WALGREENS

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

- Relieves Minor Muscle Pain for up to 8 hours

CONTAINS NO ASPIRIN

Active ingredient

(in each caplet) Acetaminophen USP, 650 mg

Purpose

Pain reliever/fever reducer

Uses

- temporarily relieves minor aches and pains due to:
- muscular aches
- backache
- minor pain of arthritis
- toothache
- premenstrual and menstrual cramps
- headache
- the common cold
- 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 and children 12 years of age and over

- 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 children under 12 years
- do not use

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.



To simulate a printed label, fold along dotted line.

HOTOTTET PARA ODS

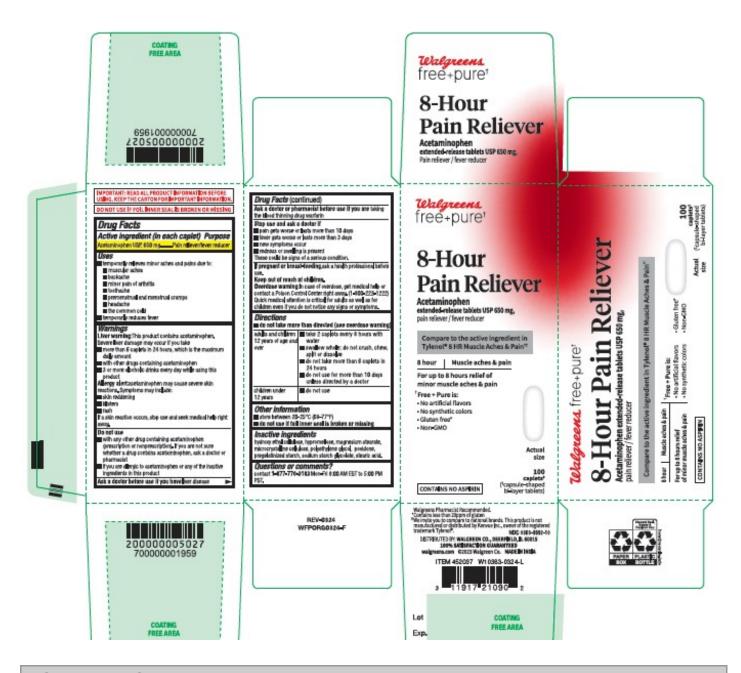


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ACETAMINOPHEN

acetaminophen tablet, film coated, extended release

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-9992
Route of Administration	ORAL		

Basis of Strength	Strength
ACETAMINOPHEN	650 mg

Inactive Ingredients	
Ingredient Name	Strength

HYPROMELLOSE 2910 (6 MPA.S) (UNII: 0WZ8WG20P6)		
HYDROXYETHYL CELLULOSE (140 CPS AT 5%) (UNII: 8136Y38GY5)		
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)		
STARCH, CORN (UNII: O8232NY3SJ)		
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)		
STEARIC ACID (UNII: 4ELV7Z65AP)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)		
POVIDONE K30 (UNII: U725QWY32X)		

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			

F	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-9992- 24	24 in 1 BOTTLE; Type 0: Not a Combination Product	09/28/2019	
2	NDC:0363-9992- 20	200 in 1 BOTTLE; Type 0: Not a Combination Product	09/28/2019	
3	NDC:0363-9992- 10	100 in 1 BOTTLE; Type 0: Not a Combination Product	09/28/2019	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA211544	09/28/2019	

Labeler - WALGREENS (008965063)

Revised: 4/2024 WALGREENS