# ACETAMINOPHEN- acetaminophen tablet, coated MEIJER DISTRIBUTION INC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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MEI-1170-2022-0328

**Drug Facts** 

#### Active ingredient (in each gelcap)

Acetaminophen 500 mg

#### **Purpose**

Pain reliever/fever reducer

#### Uses

- temporarily relieves minor aches and pains due to:
  - the common cold
  - headache
  - backache
  - minor pain of arthritis
  - toothache
  - muscular aches
  - 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 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 and children 12 years and over	<ul> <li>take 2 gelcaps every 6 hours while symptoms last</li> <li>do not take more than 6 gelcaps in 24 hours, unless directed by a doctor</li> <li>do not use for more than 10 days unless directed by a doctor</li> </ul>
children under 12 years	<ul><li>ask a doctor</li></ul>

#### Other information

- store between 20-25°C (68-77°F)
- retain carton for complete product information

## **Inactive ingredients**

colloidal silicon dioxide, croscarmellose sodium, D&C red #33, edible ink, FD&C blue #1, FD&C red #40, FD&C yellow #6, gelatin, hydroxypropyl cellulose, hypromellose, iron oxide black, iron oxide red, iron oxide yellow, polyethylene glycol, povidone, pregelatinized starch, stearic acid, titanium dioxide

## Questions or comments?

1-844-705-4384

#### PRINCIPAL DISPLAY PANEL

NDC-41250-570-03

Compare to Extra Strength Tylenol® Rapid Release Gels active ingredient† meijer®

FOR ADULTS

pain relief

500 mg Each

Acetaminophen

Pain Reliever / Fever Reducer

**EXTRA STRENGTH** 

actual size

100 GELCAPS



## **ACETAMINOPHEN**

**Product Information** 

acetaminophen tablet, coated

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Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41250-570

Route of Administration ORAL

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

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	

**ACETAMINOPHEN** 

500 mg

Inactive Ingredients	
Ingredient Name	Strength

SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GELATIN (UNII: 2G86QN327L)	
HYDROXYPROPYL CELLULOSE (1600000 WAMW) (UNII: RFW2ET671P)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX43802MRT)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E)	
STARCH, PREGELATINIZED CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	gray (red and light blue ends)	Score	no score
Shape	OVAL	Size	19mm
Flavor		Imprint Code	G1
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:41250- 570-03	1 in 1 CARTON	05/31/2022		
1		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product			
2	NDC:41250- 570-13	225 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/31/2022		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part343	05/31/2022	

## Labeler - MEIJER DISTRIBUTION INC (006959555)

Revised: 5/2022 MEIJER DISTRIBUTION INC