# PAIN RELIEF REGULAR STRENGTH- acetaminophen tablet Meijer

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-1001-2019-0903

**Drug Facts** 

## Active ingredient (in each tablet)

Acetaminophen 325 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

- adult takes more than 4,000 mg of acetaminophen in 24 hours
- child takes more than 5 doses in 24 hours, which is the maximum daily amount
- taken with other drugs containing acetaminophen
- adult has 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 the user is allergic to acetaminophen or any of the inactive ingredients in this product

**Ask a doctor before use if the user has** liver disease.

**Ask a doctor or pharmacist before use if the user is** taking the blood thinning drug warfarin

## Stop use and ask a doctor if

- pain gets worse or lasts more than 10 days in adults
- pain gets worse or lasts more than 5 days in children under 12 years
- 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 tablets every 4 to 6 hours while symptoms last</li> <li>do not take more than 10 tablets in 24 hours</li> <li>do not use for more than 10 days unless directed by a doctor</li> </ul>
children 6-11 years	<ul> <li>take 1 tablet every 4 to 6 hours while symptoms last</li> <li>do not take more than 5 tablets in 24 hours</li> <li>do not use for more than 5 days unless directed by a doctor</li> </ul>
children under 6 years	■ ask a doctor

## Other information

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

## **Inactive ingredients**

povidone, pregelatinized starch, sodium starch glycolate, stearic acid

## Questions or comments?

#### PRINCIPAL DISPLAY PANEL

NDC 41250-501-02

Compare to Regular Strength Tylenol® active ingredient†

Meijer®

Pain Relief 325 mg Each

Regular Strength

Acetaminophen | Pain Reliever | Fever Reducer

100 TABLETS



## PAIN RELIEF REGULAR STRENGTH

acetaminophen tablet

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41250-501	
Route of Administration	ORAL			

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ACETAMINO PHEN (UNII: 36209 ITL9 D) (ACETAMINO PHEN - UNII: 36209 ITL9 D)	ACETAMINOPHEN	325 mg

Inactive Ingredients		
Ingredient Name	Strength	
PO VIDO NE, UNSPECIFIED (UNII: FZ989GH94E)		
STARCH, PREGELATINIZED CORN (UNII: O8232NY3SJ)		
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)		
STEARIC ACID (UNII: 4ELV7Z65AP)		

Product Characteristics				
Color	white	Score	2 pieces	
Shape	ROUND	Size	10 mm	
Flavor		Imprint Code	M2A3;57344	
Contains				

1	Packaging			
#	t Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41250-501- 02	1 in 1 CARTON	03/01/2012	
1		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part343	03/01/2012	

## **Labeler** - Meijer (006959555)

Revised: 9/2019 Meijer