PAIN RELIEF- acetaminophen tablet 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.

Meijer Distribution, Inc. Pain Relief Drug Facts

Active ingredient (in each caplet)

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 have ever had an allergic reaction to this product or any of its ingredients

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	 take 2 caplets every 6 hours while symptoms last do not take more than 6 caplets in 24 hours, unless directed by a doctor do not use for more than 10 days unless directed by a doctor
children under 12 years	ask a doctor

Other information

• store at 20-25°C (68-77°F)

Inactive ingredients

carnauba wax, corn starch*, croscarmellose sodium*, hypromellose, polyethylene glycol, povidone, pregelatinized starch, sodium starch glycolate*, stearic acid

Questions or comments?

1-800-719-9260

^{*}may contain one or more of these ingredients

Principal Display Panel

Compare to Extra Strength Tylenol® active ingredient
EXTRA STRENGTH
pain relief 500 mg Each
Acetaminophen | Pain Reliever | Fever Reducer | For Adults
100 Caplets
actual size





Compare to ExtraStrength Tylenol*

EXTRA STRENGTH n relief 500 mg

Acetaminophen | Pain Reliever | Fever Reducer | For Adults

100 Caplets



DO NOT USE IF PRINTED SEAL UNDER IS BROKEN OR MISSING





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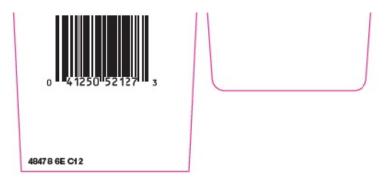
12 years

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Questions or comments? 1-800-719-9260

"This product is not manufactured or distributed by Johnson & Johnson, owner of the registered traisment Tylenol ©.



PAIN RELIEF

acetaminophen tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41250-484
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	500 mg

Inactive Ingredients			
Ingredient Name	Strength		
CARNAUBA WAX (UNII: R12CBM0EIZ)			
STARCH, CORN (UNII: O8232NY3SJ)			
HYPROMELLOSES (UNII: 3NXW29V3WO)			
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)			
PO VIDO NE (UNII: FZ989 GH94E)			
STEARIC ACID (UNII: 4ELV7Z65AP)			
CROSCARMELLOSE SODIUM (UNII: M28 OL1HH48)			

Product Characteristics				
Color	WHITE	Score	no score	
Shape	OVAL	Size	16 mm	
Flavor		Imprint Code	L484	
Contains				

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:41250-484-71	1 in 1 CARTON	08/15/1987		
1		50 in 1 BOTTLE; Type 0: Not a Combination Product			
2	NDC:41250-484-78	1 in 1 CARTON	10/28/1992		
2		100 in 1 BOTTLE; Type 0: Not a Combination Product			
3	NDC:41250-484-90	500 in 1 BOTTLE; Type 0: Not a Combination Product	03/13/1992		

4	NDC:41250-484-76	1 in 1 CARTON	03/20/1992	
4		120 in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:41250-484-02	550 in 1 BOTTLE; Type 0: Not a Combination Product	06/28/2013	04/06/2016
6	NDC:41250-484-82	2 in 1 PACKAGE	10/28/1992	
6		1 in 1 CARTON		
6		100 in 1 BOTTLE; Type 0: Not a Combination Product		
7	NDC:41250-484-62	1 in 1 CARTON	06/16/2017	
7		24 in 1 BOTTLE; 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	08/15/1987	

Labeler - Meijer Distribution Inc (006959555)

Revised: 1/2019 Meijer Distribution Inc