

CAREALL NON ASPIRIN EXTRA STRENGTH- acetaminophen tablet
New World Imports, Inc

Drug Facts

Acetaminophen 500 mg

Pain reliever / Fever reducer

In case of accidental overdose, contact a doctor or Poison Control Center (1-800-222-1222) right away. Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

For the temporary relief of minor aches and pains associated with: the common cold, headache, toothache, muscular aches, backache, minor pain from arthritis, premenstrual and menstrual cramps

Temporarily reduces fever

Liver Warning:This product contains acetaminophen. Severe liver damage may occur if you take:

- more than 4000 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 for more than 10 days
- Fever gets worse or lasts for 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.

Do not exceed recommended dosage.

Do not use more than directed (see overdose warnings)


Adults and children 12 years of age and older:

- 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 more than 10 days unless directed by a doctor.

Children under 12 years of age: consult a doctor

Corn Starch, povidone, pregelatinized starch, sodium starch glycolate, stearic acid

NDC 51824-049-01




Non-Aspirin


Extra Strength

▶ Pain Reliever
▶ Fever Reducer

See new warnings information



Compare to the active ingredient of Extra Strength **TYLENOL**®



100 Acetaminophen Caplets | 500mg each

Drug Facts

Active ingredient (in each caplet) *Purposes*
Acetaminophen 500 mgPain reliever/Fever reducer

Uses temporary relief of minor aches and pains associated with ■ common cold ■ headache ■ toothache ■ muscular aches ■ backache ■ arthritis ■ menstrual cramps ■ and reduction of fever

Warnings
Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take
 ■ more than 4000 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
 ■ if you are allergic to acetaminophen ■ with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.

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

Drug Facts (continue under label)

*This product is not manufactured or distributed by McNeil Consumer Healthcare, owner of the registered trademark Tylenol®. Distributed by: INWI, Inc., 160 Athens Way, Nashville, TN 37228



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Lot Exp.

PEEL HERE

Drug Facts (Continued)

Stop use and ask a doctor if ■ symptoms do not improve ■ pain gets worse or lasts for more than 10 days ■ fever gets worse or lasts for more than 3 days ■ new symptoms occur ■ redness or swelling is present ■ a rare sensitivity reaction occurs

You may report side effects to 1-888-952-0050
If pregnant or breast-feeding, ask a health professional before use

Keep out of reach of children. In case of accidental overdose, contact a doctor or Poison Control Center immediately. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.
Do not exceed recommended dosage.

Directions
 ■ do not use more than directed (see overdose warning)
 ■ adults and children 12 years and over: take 2 caplets every 6 hours. Do not take more than 8 caplets in 24 hours. ■ Do not use more than 10 days unless directed by a doctor
 ■ children under 12 years: do not use this product in children under 12 years of age; this will provide more than the recommended dose (overdose) and may cause liver damage

Overdose warning: Taking more than the recommended dose (overdose) may cause liver damage. In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away. Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Other information
 ■ Store between 20°-25°C (68°-77°F) ■ do not use if imprinted safety seal under cap is broken or missing

Inactive ingredients
 corn starch, pregelatinized starch, stearic acid. May contain povidone and sodium starch glycolate.

CAREALL NON ASPIRIN EXTRA STRENGTH

acetaminophen tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51824-049
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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Acetaminophen (UNII: 36209ITL9D) (Acetaminophen - UNII:36209ITL9D)	Acetaminophen	500 mg
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Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
Propylene Glycol (UNII: 6DC9Q167V3)	
Stearic Acid (UNII: 4ELV7Z65AP)	
Povidone (UNII: FZ989GH94E)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	

Product Characteristics

Color	white (White)	Score	no score
Shape	OVAL (Oval)	Size	18mm
Flavor		Imprint Code	AZ;328
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51824-049-24	24 in 1 CASE	09/17/2015	
1	NDC:51824-049-01	100 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 Drug	M013	09/17/2015	

Labeler - New World Imports, Inc (075372276)

Revised: 11/2023

New World Imports, Inc