

PHARBETOL- acetaminophen tablet
Pharbest Pharmaceuticals, Inc.

Pharbetol Extra Strength Tablet

Active ingredient (in each tablet)

Acetaminophen 500 mg

Purpose

Pain reliever/fever reducer

Uses

- temporarily relieves minor aches and pains due to:
- 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 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 more than 10 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur

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 don't notice any signs or symptoms.

Directions

- **do not take more than directed (see overdose warning).**

adult and children 12 years and over	<ul style="list-style-type: none">• take 2 tablets every 6 hours while symptoms last• do not take more than 6 tablets 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 20-25⁰C (68-77⁰F)

Inactive ingredients

povidone, pregelatinized corn starch, sodium starch glycolate, stearic acid

Questions or comments?

(866) 562-2756 (Mon - Fri 8 AM to 4 PM EST)

PHARBEST

NDC 16103-376-11

manufactured in the USA

Extra Strength Contains no Aspirin

***Compare to the active ingredient in Extra Strength Tylenol® Caplet**

PHARBETOL

Acetaminophen 500 mg each

Pain Reliever/Fever Reducer

THIS PACKAGE FOR HOUSEHOLD WITHOUT YOUNG CHILDREN

1000 TABLETS

TAMPER EVIDENT: DO NOT USE IF SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

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Drug Facts
Active ingredient (in each tablet)
Acetaminophen 500 mg. Pain reliever/fever reducer.

Uses
temporarily relieves minor aches and pains due to: 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, or 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 a 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, redness or swelling is present, new symptoms occur. These could be signs of a serious condition.

If pregnant or live ask-is editing, 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 don't notice any signs or symptoms.

Directions • do not take more than directed (see overdose warning)
adults and children 12 years and over: take 2 tablets every 6 hours while symptoms last. Do not take more than 8 tablets in 24 hours, unless directed by a doctor.
children under 12 years: do not use for more than 10 days unless directed by a doctor. ask a doctor.

Other information
store at 20-25°C (68-77°F).

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

Questions or comments? (866) 562-2756 (Mon-Fri 8 AM to 4 PM EST)

Manufactured by:
Pharbest Pharmaceuticals, Inc.
Farmingdale, NY 11735

Org. 08/21

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PHARBETOL

acetaminophen tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:16103-376
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg

Inactive Ingredients

Ingredient Name	Strength
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POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
STARCH, CORN (UNII: O8232NY3SJ)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	12mm
Flavor		Imprint Code	PH044
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:16103-376-06	1 in 1 CARTON	01/10/2006	
1		50 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:16103-376-08	1 in 1 CARTON	01/10/2006	
2		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:16103-376-11	1000 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/10/2006	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	01/10/2006	

Labeler - Pharbest Pharmaceuticals, Inc. (557054835)

Establishment

Name	Address	ID/FEI	Business Operations
Pharbest Pharmaceuticals, Inc.		557054835	analysis(16103-376) , manufacture(16103-376) , pack(16103-376) , label(16103-376)

Revised: 12/2025

Pharbest Pharmaceuticals, Inc.