

PHARBETOL- acetaminophen tablet
Pharbest Pharmaceuticals, Inc.

Pharbetol Extra Strength Tablet

Drug Facts

Active ingredient (in each tablet)

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 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 4 to 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 between 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 500mg

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 or 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 redness, 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 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)
adults and children 12 years and over take 2 tablets every 6 hours while symptoms last
do not take more than 6 tablets in 24 hours, unless directed by a doctor
years and over 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 povidone, pregelatinized corn starch, sodium starch glycolate, stearic acid

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

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

*This product is not manufactured or distributed by McNeil Consumer Healthcare Division of McNeil-PPC, Inc., owner of the registered trademark "Tylenol".

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
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	M	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: 1/2024

Pharbest Pharmaceuticals, Inc.