#### PHARBETOL- acetaminophen tablet Pharbest Pharmaceuticals, Inc.

-----

Pharbetol Extra Stength 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> <li>take 2 tablets, every 4 to 6 hours while symptoms last</li> <li>do not take more than 6 tablets in 24 hours, unless directed by a doctor</li> <li>do not use for more than 10 days unless directed by a doctor</li> </ul>
children under 12 years	Ask a doctor

# Other information

• store between 20-25<sup>0</sup>C (68-77<sup>0</sup>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)

#### PHARBEST

NDC 16103-376-11

manufactured in the USA

Extra Strength Contains no Aspirin

\*Compare to the active ingredient in Extra Strength Tylenol<sup>®</sup> Caplet

#### PHARBETOL

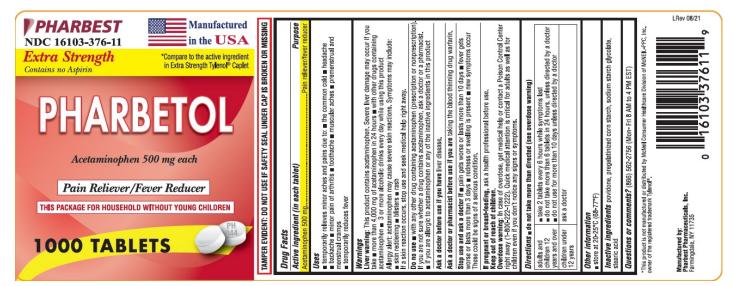
Acetaminophen 500mg

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



PHARBETOL					
acetaminophen tablet					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC		NDC:16103-376	
Route of Administration	ORAL				
Active Ingredient/Active Moiety					
Ingre	Basis of St	trength	Strength		
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII: 36209ITL9D)			ACETAMINOPH	EN	500 mg

Inactive Ingredients				
Ingredient Name	Strength			
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)				
STARCH, CORN (UNII: 08232NY3SJ)				
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

#	ltem 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	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
OTC Monograph Drug	Μ	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.