PHARBETOL- acetaminophen tablet
Proficient Rx LP

Drug Facts

Active ingredient (in each tablet)

Acetaminophen 500mg

Purpose

Pain reliever/fever reducer

Uses

Temporarily relieves minor aches and pains due to:

- headache
- backache
- minor pain of arthritis
- toothache
- muscular aches
- menstrual cramps

Temporarily reduces fever

Warnings

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

- more than 8 tablets in 24 hours, which is the maximum daily amount
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

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.
- 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:

Taking more than the recommended dose (overdose) may cause liver damage. In the case of overdose, get medical help or contact a Poison Control Center right away.

Directions

- **do not take more than directed (see overdose warning).**
- **Adult and Children 12 years and over:** take 2 tablets, every 4 to 6 hours while symptoms last. Do not take more than 8 tablets in 24 hours. Do not take for more than 10 days unless directed by a doctor.
- **Children under 12 years:** do not use adult extra strength product in children under 12 years of age; this will provide more than recommended dose (overdose) and may cause liver damage.

Inactive ingredients

Povidone, Pregelatinized corn starch, sodium starch glycolate, stearic acid.

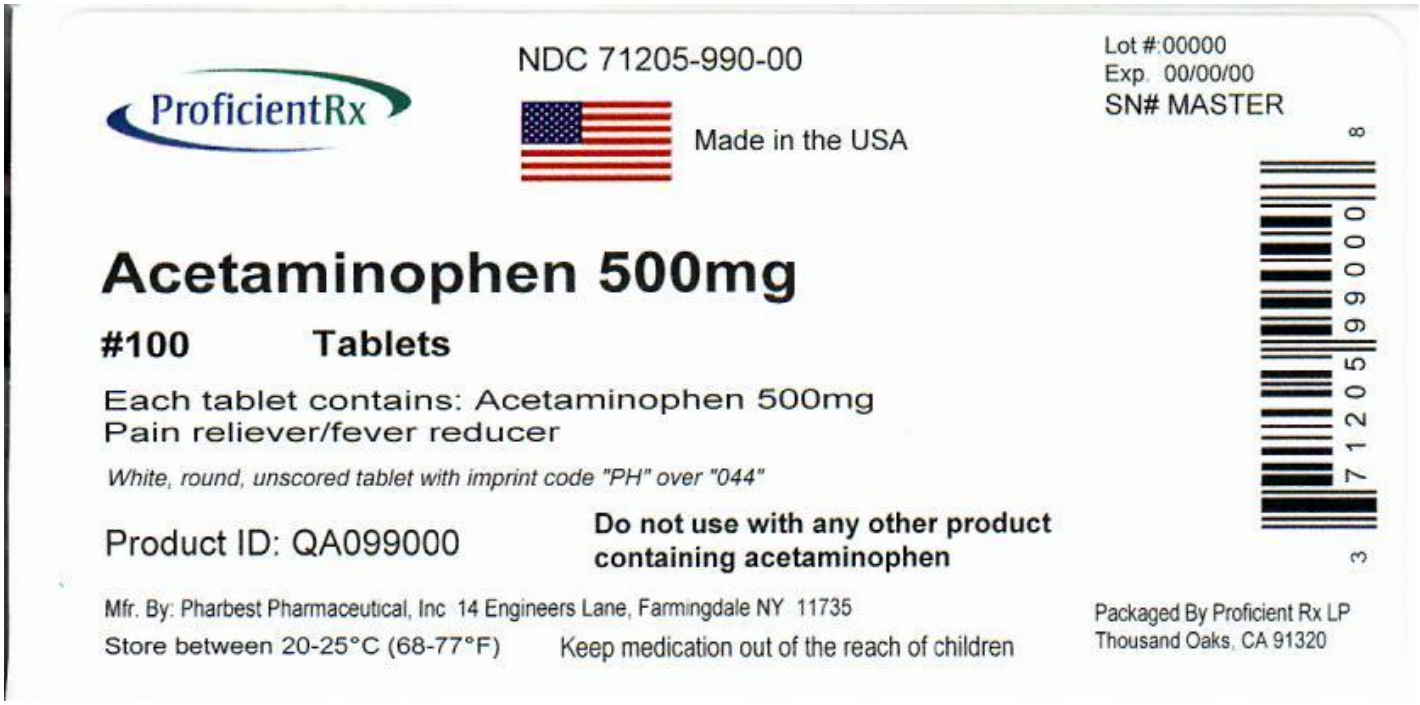
Questions?

Adverse drug event call: (866) 562-2756

Repackaged by:

Proficient Rx LP

Thousand Oaks, CA 91320



NDC 71205-990-00

Manufactured in the USA

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

Do not use any other product containing acetaminophen

Acetaminophen 500mg

Pain Reliever Fever Reducer

100 Tablets

PHARBETOL

acetaminophen tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71205-990(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 K30 (UNII: U725QWY32X)	
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:71205-990-50	50 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/01/2020	
2	NDC:71205-990-00	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/01/2020	
3	NDC:71205-990-55	500 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/01/2020	
4	NDC:71205-990-11	1000 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/01/2020	

Marketing Information

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

Labeler - Proficient Rx LP (079196022)

Establishment

Name	Address	ID/FEI	Business Operations
Proficient Rx LP		079196022	REPACK(71205-990) , RELABEL(71205-990)