PHARBETOL- acetaminophen tablet
Proficient Rx LP

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 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

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
- 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: In case of overdose, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

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 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

- **Tamper Evident: do not use if imprinted safety seal under cap is broken or missing**
- store between 20-25^{0}C (68-77^{0}F)

Inactive ingredients

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

Questions?

Adverse drug event call: 1-866-562-2756 Mon - Fri: 8 AM to 4 PM

Repackaged By:

Proficient Rx LP

Thousand Oaks, CA 91320

NDC 71205-478-50

manufactured in the USA

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

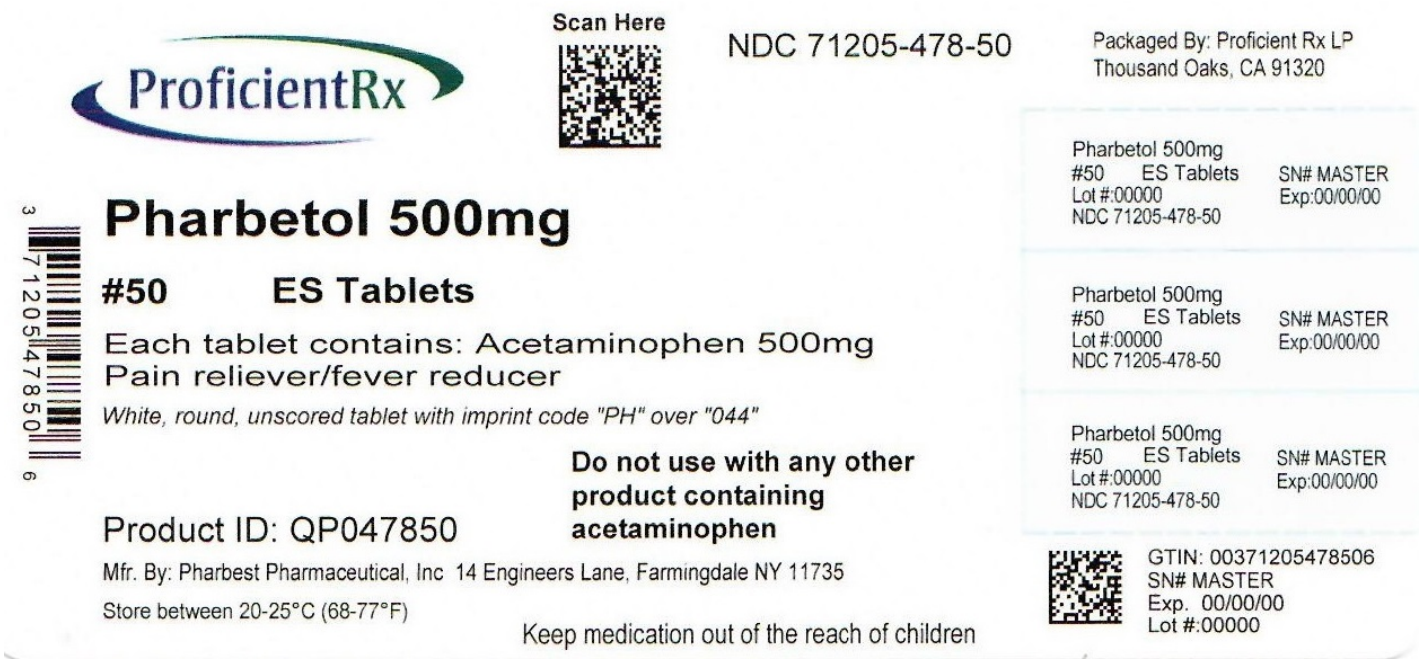
Do not use with any other product containing acetaminophen

PHARBETOL

Acetaminophen 500mg

Pain Reliever • Fever Reducer

50 TABLETS



PHARBETOL

acetaminophen tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71205-478(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:71205-478-20	20 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/26/2024	
2	NDC:71205-478-24	24 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/30/2020	
3	NDC:71205-478-30	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/30/2020	
4	NDC:71205-478-40	40 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/12/2020	
5	NDC:71205-478-50	50 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/12/2020	
6	NDC:71205-478-60	60 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/30/2020	
7	NDC:71205-478-90	90 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/30/2020	

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(71205-478) , RELABEL(71205-478)

Revised: 1/2024

Proficient Rx LP