

**PHARBETOL- acetaminophen tablet**  
**ATLANTIC BIOLOGICALS CORP.**

*Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.*

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**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	<ul style="list-style-type: none"><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**

- **Tamper Evident: do not use if imprinted safety seal under cap is broken or missing**
- 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?**

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

**DISTRIBUTED BY:**

ATLANTIC BIOLOGICALS CORP.

MIAMI, FL 33179

**NDC 17856-0376-01**

**PHARBETOL**

**Acetaminophen 500mg**

**Pain Reliever • Fever Reducer**

**100 TABLETS**

**17856-0376-01**  
**PHARBETOL**  
**(ACETAMINOPHEN)**  
**EXTRA STRENGTH**  
**500 MG TABLETS**



See package insert for indications and dosage schedule

Store at 20°-25°C (68°-77°F ) PAIN RELIEVER & FEVER REDUCER  
 \*\* Keep this and all medication out of the reach of children\*\*



17856-0376-01

Dosage 500 MG TABLETS

**PHARBETOL**  
**(ACETAMINOPHEN)**

Qty: 100 TABLETS



GTIN: 00117856037613

S/N: 01840001

Exp: 03/07/23

Lot: 018400



Packaged by: Unit Dose Solutions  
 Morrisville, NC 27560

Distributed by: Atlantic Biologicals Corp.  
 Miami FL 33179

Rev.08/21

**Call to Reorder: 800.509.7592**

**PHARBETOL**

acetaminophen tablet

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:17856-0376(NDC:16103-376)
<b>Route of Administration</b>	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:17856-0376-1	100 in 1 BOX, UNIT-DOSE	09/08/2022	
1	NDC:17856-0376-2	1 in 1 POUCH; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part343	01/10/2006	

**Labeler** - ATLANTIC BIOLOGICALS CORP. (047437707)

**Registrant** - ATLANTIC BIOLOGICALS CORP. (047437707)

### Establishment

Name	Address	ID/FEI	Business Operations
UNIT DOSE SOLUTIONS		360804194	repack(17856-0376)

Revised: 9/2022

ATLANTIC BIOLOGICALS CORP.