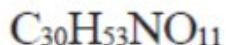
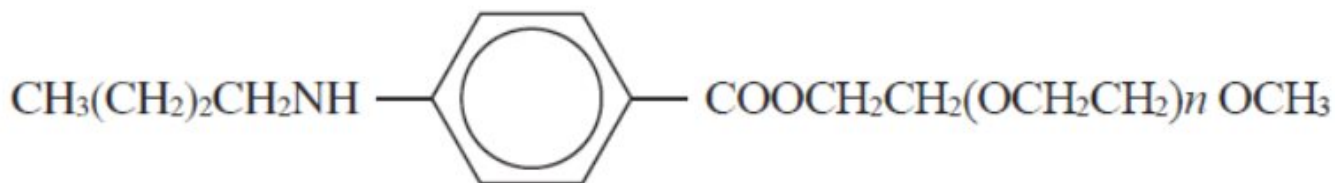


**BENZONATATE- benzonatate capsule**  
**PD-Rx Pharmaceuticals, Inc.**

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**Benzonatate Capsules, USP**

**DESCRIPTION**

Benzonatate capsules, USP, a non-narcotic oral antitussive agent, is 2, 5, 8, 11, 14, 17, 20, 23, 26- onaoxaoctacosan-28-ylp-(butylamino) benzoate; with a molecular weight of 603.7.



Each benzonatate capsules, USP for oral administration contains 100 mg or 200 mg benzonatate, USP. In addition, each capsule contains the following inactive ingredients: D&C Yellow #10, gelatin, glycerin, lecithin, light mineral oil, propylene glycol, purified water, shellac glaze, titanium dioxide, and white edible ink.

**CLINICAL PHARMACOLOGY**

Benzonatate capsules act peripherally by anesthetizing the stretch receptors located in the respiratory passages, lungs, and pleura by dampening their activity and thereby reducing the cough reflex at its source. It begins to act within 15 to 20 minutes and its effect lasts for 3 to 8 hours. Benzonatate capsules have no inhibitory effect on the respiratory center in recommended dosage.

**INDICATIONS AND USAGE**

Benzonatate capsules, USP are indicated for the symptomatic relief of cough.

**CONTRAINDICATIONS**

Hypersensitivity to benzonatate or related compounds.

**WARNINGS**

**Hypersensitivity**

Severe hypersensitivity reactions (including bronchospasm, laryngospasm and cardiovascular collapse) have been reported which are possibly related to local anesthesia from sucking or chewing the capsule instead of swallowing it. Severe

reactions have required intervention with vasopressor agents and supportive measures.

### **Psychiatric Effects**

Isolated instances of bizarre behavior, including mental confusion and visual hallucinations, have also been reported in patients taking Benzonatate capsules in combination with other prescribed drugs.

### **Accidental Ingestion and Death in Children**

Keep benzonatate capsules out of reach of children. Accidental ingestion of benzonatate capsules resulting in death has been reported in children below age 10. Signs and symptoms of overdose have been reported within 15-20 minutes and death has been reported within one hour of ingestion. If accidental ingestion occurs, seek medical attention immediately (see **OVERDOSAGE**).

### **PRECAUTIONS**

Benzonatate is chemically related to anesthetic agents of the para-amino-benzoic acid class (e.g. procaine; tetracaine) and has been associated with adverse CNS effects possibly related to a prior sensitivity to related agents or interaction with concomitant medication.

#### **Information for Patients:**

Swallow benzonatate capsules whole. Do not break, chew, dissolve, cut, or crush benzonatate capsules. Release of benzonatate from the capsule in the mouth can produce a temporary local anesthesia of the oral mucosa and choking could occur. If numbness or tingling of the tongue, mouth, throat, or face occurs, refrain from oral ingestion of food or liquids until the numbness has resolved. If the symptoms worsen or persist, seek medical attention.

Keep benzonatate capsules out of reach of children. Accidental ingestion resulting in death has been reported in children. Signs and symptoms of overdose have been reported within 15-20 minutes and death has been reported within one hour of ingestion. Signs and symptoms may include restlessness, tremors, convulsions, coma and cardiac arrest. If accidental ingestion occurs, seek medical attention immediately.

Overdosage resulting in death may occur in adults.

Do not exceed a single dose of 200 mg and a total daily dosage of 600 mg. If you miss a dose of benzonatate capsules, skip that dose and take the next dose at the next scheduled time. Do not take 2 doses of benzonatate capsules at one time.

#### **Usage in Pregnancy**

##### **PREGNANCY CATEGORY C**

Animal reproduction studies have not been conducted with benzonatate capsules. It is also not known whether benzonatate capsules can cause fetal harm when administered to a pregnant woman or can affect reproduction capacity. Benzonatate capsules should be given to a pregnant woman only if clearly needed.

## **Nursing Mothers**

It is not known whether this drug is excreted in human milk. Because many drugs are excreted in human milk caution should be exercised when benzonatate capsules are administered to a nursing woman.

## **Carcinogenesis, Mutagenesis, Impairment of Fertility**

Carcinogenicity, mutagenicity, and reproduction studies have not been conducted with benzonatate capsules..

## **Pediatric Use**

Safety and effectiveness in children below the age of 10 have not been established. Accidental ingestion resulting in death has been reported in children below age 10. Keep out of reach of children.

## **ADVERSE REACTIONS**

Potential Adverse Reactions to benzonatate capsules may include:

**Hypersensitivity reactions** including bronchospasm, laryngospasm, cardiovascular collapse possibly related to local anesthesia from chewing or sucking the capsule.

### **CNS**

sedation; headache; dizziness; mental confusion; visual hallucinations.

### **GI**

constipation; nausea; GI upset.

### **Dermatologic**

pruritus; skin eruptions.

### **Other**

nasal congestion; sensation of burning in the eyes; vague "chilly" sensation; numbness of the chest; hypersensitivity.

Deliberate or accidental overdose has resulted in death, particularly in children.

### **Hypersensitivity reactions**

including bronchospasm, laryngospasm, cardiovascular collapse possibly related to local anesthesia from chewing or sucking the capsule.

## **OVERDOSAGE**

Intentional and unintentional overdose may result in death, particularly in children.

The drug is chemically related to tetracaine and other topical anesthetics and shares various aspects of their pharmacology and toxicology. Drugs of this type are generally well absorbed after ingestion.

## Signs and Symptoms

The signs and symptoms of overdose of benzonatate capsules have been reported within 15-20 minutes. If capsules are chewed or dissolved in the mouth, oropharyngeal anesthesia will develop rapidly, which may cause choking and airway compromise.

CNS stimulation may cause restlessness and tremors which may proceed to clonic convulsions followed by profound CNS depression. Convulsions, coma, cerebral edema and cardiac arrest leading to death have been reported within 1 hour of ingestion.

## Treatment

In case of overdose, seek medical attention immediately. Evacuate gastric contents and administer copious amounts of activated charcoal slurry. Even in the conscious patient, cough and gag reflexes may be so depressed as to necessitate special attention to protection against aspiration of gastric contents and orally administered materials. Convulsions should be treated with a short-acting barbiturate given intravenously and carefully titrated for the smallest effective dosage. Intensive support of respiration and cardiovascular-renal function is an essential feature of the treatment of severe intoxication from overdosage.

Do not use CNS stimulants.

## DOSAGE AND ADMINISTRATION

Adults and Children over 10 years of age: Usual dose is one 100 mg or 200 mg capsule three times a day as needed for cough. If necessary to control cough, up to 600 mg daily in three divided doses may be given. **Benzonatate capsules should be swallowed whole.** Benzonatate capsules are not to be broken, chewed, dissolved, cut or crushed.

## HOW SUPPLIED

Benzonatate Capsules USP, 100 mg are clear yellow, oval-shaped softgel capsules imprinted with "PC14" and are supplied as follows:

NDC 72789-230-06 in bottles of 6 capsules

NDC 72789-230-12 in bottles of 12 capsules

NDC 72789-230-15 in bottles of 15 capsules

NDC 72789-230-20 in bottles of 20 capsules

NDC 72789-230-21 in bottles of 21 capsules

NDC 72789-230-30 in bottles of 30 capsules

Store at 20° to 25° C (68° to 77° F) [see USP Controlled Room Temperature].

PROTECT FROM LIGHT

Dispense in tight (USP), child-resistant containers.

**Rx Only**

# PACKAGE/LABEL PRINCIPAL DISPLAY PANEL - 100 mg

DO NOT CRUSH OR CHEW CAPSULES.

**R only** WARNING: KEEP THIS OUT OF THE REACH OF CHILDREN  
DOSAGE and STORAGE: SEE PACKAGE INSERT

72789-230-20	72789-230-20	72789-230-20
BENZONATATE USP	BENZONATATE USP	BENZONATATE USP
100 MG	100 MG	100 MG
20 CAPSULES	20 CAPSULES	20 CAPSULES
ReOrder # 111285	ReOrder # 111285	ReOrder # 111285
LOT C23D66	LOT C23D66	LOT C23D66
EXP 08/2025	EXP 08/2025	EXP 08/2025

CALL YOUR DOCTOR FOR MEDICAL ADVICE ABOUT SIDE EFFECTS.  
YOU MAY REPORT SIDE EFFECTS TO THE FDA AT 1-800-FDA-1088

TAKE \_\_\_\_\_ CAPSULE(S) \_\_\_\_\_ TIMES A DAY.  
TOME \_\_\_\_\_ CAPSULA(S) \_\_\_\_\_ VECES AL DIA.

Each CAPSULE Contains: BENZONATATE USP 100 MG

NDC: 72789-230-20



**BENZONATATE USP**

**100 MG**

**20 CAPSULES**



GTIN: 00372789230207  
SNO: C23D66000001  
EXP: 08/2025  
LOT: C23D66

YELLOW  
ORGANOLEPTIC MARKINGS:  
PC 14  
CAPSULE

372799230207  
42806071405  
HUMANWELL PHARACAP PHARM.  
WUHAN, HUBEI 430206  
CHINA

## BENZONATATE

benzonatate capsule

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:72789-230(NDC:42806-714)
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>BENZONATATE</b> (UNII: 5P4DHS6ENR) (BENZONATATE - UNII:5P4DHS6ENR)	BENZONATATE	100 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>D&amp;C YELLOW NO. 10</b> (UNII: 35SW5USQ3G)	
<b>GELATIN</b> (UNII: 2G86QN327L)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>LECITHIN, SOYBEAN</b> (UNII: 1DI56QDM62)	
<b>LIGHT MINERAL OIL</b> (UNII: N6K5787QVP)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>SHELLAC</b> (UNII: 46N107B710)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>WATER</b> (UNII: 059QF0KO0R)	

## Product Characteristics

<b>Color</b>	yellow	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	7mm
<b>Flavor</b>		<b>Imprint Code</b>	PC14
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72789-230-15	15 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/02/2022	
2	NDC:72789-230-21	21 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/02/2022	
3	NDC:72789-230-30	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/02/2022	
4	NDC:72789-230-20	20 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/16/2022	
5	NDC:72789-230-12	12 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/21/2022	
6	NDC:72789-230-06	6 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/16/2022	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA206948	12/21/2018	

**Labeler** - PD-Rx Pharmaceuticals, Inc. (156893695)

**Registrant** - PD-Rx Pharmaceuticals, Inc. (156893695)

## Establishment

Name	Address	ID/FEI	Business Operations
PD-Rx Pharmaceuticals, Inc.		156893695	repack(72789-230)

Revised: 4/2023

PD-Rx Pharmaceuticals, Inc.