

**BENAZEPRIL HYDROCHLORIDE- benazepril hydrochloride tablet, film coated**  
**Northwind Pharmaceuticals**

NDC: 51655-067-52

MFG: 63304-0339-05

Benazepril Hydrochloride 20 MG

30 Tablets

Rx only

Lot# NW89620001

Exp Date: 2/2015

Each tablet contains: Benasepril hydrochloride, USP....20 MG

Dosage: See package outsert for full prescribing information

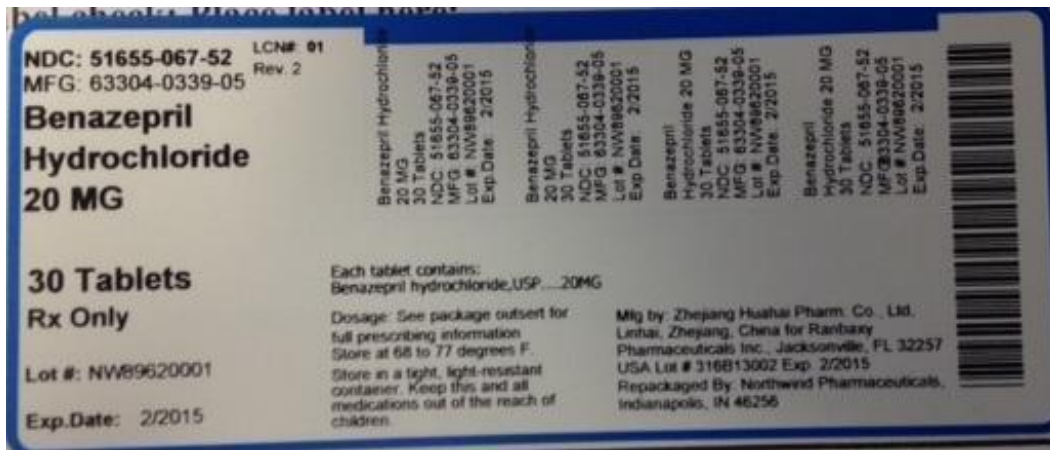
Store at 60 to 77 degrees F.

Store in a tight, light-resistanct container. Keep this and all medications out the the reach of children.

Mfg by: Zhejiang Huahai Pharm. Co., Ltd Linhai, Zheijang, China for Ranbaxy Pharmaceuticals Inc, Jacksonville, FL 32257 USA

lot # 316B13002 Exp 2/2015

Repackaged by Northwind Pharmaceuticals, Indianapolis, IN 46256



**BENAZEPRIL HYDROCHLORIDE**

benazepril hydrochloride tablet, film coated

**Product Information**

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:51655-067(NDC:63304-339)
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
-----------------	-------------------	----------

BENAZEPRIL HYDROCHLORIDE (UNII: N1SN99T69T) (BENAZEPRILAT - UNII:JRM708L703)	BENAZEPRIL HYDROCHLORIDE	20 mg
--	--------------------------	-------

### Product Characteristics

<b>Color</b>	gray	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	5mm
<b>Flavor</b>		<b>Imprint Code</b>	343
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51655-067-52	30 in 1 BOTTLE, DISPENSING		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076118	05/30/2014	

**Labeler** - Northwind Pharmaceuticals (036986393)

**Registrant** - Northwind Pharmaceuticals (036986393)

### Establishment

Name	Address	ID/FEI	Business Operations
Northwind Pharmaceuticals		036986393	repack(51655-067)

Revised: 6/2014

Northwind Pharmaceuticals