

FENOFIBRATE- fenofibrate tablet
Northwind Pharmaceuticals

NDC: 51655-009-52

MFG: 0115-552-10

Fenofibrate 160mg

30 Tablets

Rx Only

Lot# NW89560001

Exp Date: 09/2015

Each tablet contains: 160mg of fenofibrate

Dosage: See package insert

Store at 60-77 degrees F. Store in a tight, light-resistant container (See USP)

Keep out of the reach of children.

Dist. By: Global Pharmaceuticals

Division of IMPAX Laboratories, Inc

Philadelphia, PA 19124 USA

Product of Taiwan Lot: 10008621

Repackaged by Northwind Pharmaceuticals, Indianapolis, IN 46256



FENOFIBRATE

fenofibrate tablet

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:51655-009(NDC:0115-5522)
Route of Administration	oral		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FENOFIBRATE (UNII: U202363UOS) (FENOFIBRATE - UNII:U202363UOS)	FENOFIBRATE	160 mg in 30

Product Characteristics

Color	white	Score	no score
Shape	OVAL	Size	15mm
Flavor		Imprint Code	G;352
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51655-009-52	30 in 1 BOTTLE, DISPENSING		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076509	05/07/2014	

Labeler - Northwind Pharmaceuticals (036986393)**Registrant** - Northwind Pharmaceuticals (036986393)**Establishment**

Name	Address	ID/FEI	Business Operations
Northwind Pharmaceuticals		036986393	repack(51655-009)

Revised: 5/2014

Northwind Pharmaceuticals