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Drug Details

Drug Name(s)	FENOFIBRIC ACID
FDA Application No.	(ANDA) 200920
Active Ingredient(s)	CHOLINE FENOFIBRATE
Company	ACTAVIS ELIZABETH
Original Approval or Tentative Approval Date	October 7, 2015

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- **Labels are not available**

Products on Application (ANDA) #200920

Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
FENOFIBRIC ACID	CHOLINE FENOFIBRATE	EQ 45MG FENOFIBRIC ACID	CAPSULE, DELAYED RELEASE;ORAL	Prescription No	AB
FENOFIBRIC ACID	CHOLINE FENOFIBRATE	EQ 135MG FENOFIBRIC ACID	CAPSULE, DELAYED RELEASE;ORAL	Prescription No	AB

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