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Drug Details

Drug Name(s)	FENOFIBRATE
FDA Application No.	(ANDA) 204598
Active Ingredient(s)	FENOFIBRATE
Company	HETERO LABS LTD III
Original Approval or Tentative Approval Date	July 12, 2016

- [Therapeutic Equivalents](#)
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- **Labels are not available**

Products on Application (ANDA) #204598

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
FENOFIBRATE	FENOFIBRATE	48MG	TABLET;ORAL	Prescription	No	AB
FENOFIBRATE	FENOFIBRATE	145MG	TABLET;ORAL	Prescription	No	AB

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