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Drug Details

Drug Name(s)	FELODIPINE
FDA Application No.	(ANDA) 203032
Active Ingredient(s)	FELODIPINE
Company	ORCHID HLTHCARE
Original Approval or Tentative Approval Date	May 21, 2015

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

Products on Application (ANDA) #203032

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
FELODIPINE	FELODIPINE	2.5MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB
FELODIPINE	FELODIPINE	5MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB
FELODIPINE	FELODIPINE	10MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB

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