Drug Details

Drug Name(s)	**IRBESARTAN**
FDA Application No.	**(ANDA) 203685**
Active Ingredient(s)	**IRBESARTAN**
Company	**AJANTA PHARMA LTD**
Original Approval or Tentative Approval Date	**December 10, 2015**

- **There are no Therapeutic Equivalents**
- **Labels are not available**
- **Approval History, Letters, Reviews, and Related Documents**

Products on Application (ANDA) #203685
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
IRBESARTAN	IRBESARTAN	75MG	TABLET;ORAL	None (Tentative Approval)	No	None
IRBESARTAN	IRBESARTAN	150MG	TABLET;ORAL	None (Tentative Approval)	No	None
IRBESARTAN	IRBESARTAN	300MG	TABLET;ORAL	None (Tentative Approval)	No	None

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