Drug Details

Drug Name(s)	**OLANZAPINE**
FDA Application No.	**(ANDA) 206711**
Active Ingredient(s)	**OLANZAPINE**
Company	**AJANTA PHARMA LTD**
Original Approval or Tentative Approval Date	**August 30, 2016**

- **Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Labels are not available**

Products on Application (ANDA) #206711
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
OLANZAPINE	OLANZAPINE	2.5MG	TABLET;ORAL	Prescription	No	AB
OLANZAPINE	OLANZAPINE	5MG	TABLET;ORAL	Prescription	No	AB
OLANZAPINE	OLANZAPINE	7.5MG	TABLET;ORAL	Prescription	No	AB
OLANZAPINE	OLANZAPINE	10MG	TABLET;ORAL	Prescription	No	AB
OLANZAPINE	OLANZAPINE	15MG	TABLET;ORAL	Prescription	No	AB
OLANZAPINE	OLANZAPINE	20MG	TABLET;ORAL	Prescription	No	AB

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