Drugs@FDA
FDA Approved Drug Products

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Drug Details

Drug Name(s)	**ZYTIGA**
FDA Application No.	**(NDA) 202379**
Active Ingredient(s)	**ABIRATERONE ACETATE**
Company	**JANSSEN BIOTECH**
Original Approval or Tentative Approval Date	**April 28, 2011**
Chemical Type	**1 New molecular entity (NME)**
Review Classification	**P Priority review drug**

- **There are no Therapeutic Equivalents**
- **Label Information**
- **Approval History, Letters, Reviews, and Related Documents**

Products on Application (NDA) #202379
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
ZYTIGA	ABIRATERONE ACETATE	250MG	TABLET;ORAL	Prescription	Yes	None

Back to Top | Back to Previous Page | Back to Drugs@FDA Home

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