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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](#)
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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

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