Drug Details

Drug Name(s)	**OXALIPLATIN**
FDA Application No.	**(ANDA) 204616**
Active Ingredient(s)	**OXALIPLATIN**
Company	**QILU PHARM CO LTD**
Original Approval or Tentative Approval Date	**May 11, 2016**

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

Products on Application (ANDA) #204616
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
OXALIPLATIN	OXALIPLATIN	50MG/VIAL	INJECTABLE;IV (INFUSION)	Prescription	No	AP
OXALIPLATIN	OXALIPLATIN	100MG/VIAL	INJECTABLE;IV (INFUSION)	Prescription	No	AP

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