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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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