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Drug Details

Drug Name(s)	OXYCODONE HYDROCHLORIDE
FDA Application No.	(ANDA) 206456
Active Ingredient(s)	OXYCODONE HYDROCHLORIDE
Company	WOCKHARDT BIO AG
Original Approval or Tentative Approval Date	June 17, 2015

- [Therapeutic Equivalents](#)
- [Approval History, Letters, Reviews, and Related Documents](#)
- **Labels are not available**

Products on Application (ANDA) #206456

Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength Dosage Form/Route	Marketing Status	RLD TE Code
OXYCODONE HYDROCHLORIDE	OXYCODONE HYDROCHLORIDE	5MG/5ML SOLUTION;ORAL	Prescription	No AA

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