Drugs@FDA
FDA Approved Drug Products

Drug Details

Drug Name(s)	**OXYMORPHONE HYDROCHLORIDE**
FDA Application No.	**(ANDA) 204459**
Active Ingredient(s)	**OXYMORPHONE HYDROCHLORIDE**
Company	**AUROLIFE PHARMA LLC**
Original Approval or Tentative Approval Date	**April 26, 2016**

- **Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Labels are not available**
- **REMS[11]**

Products on Application (ANDA) #204459
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
OXYMORPHONE HYDROCHLORIDE	OXYMORPHONE HYDROCHLORIDE	5MG	TABLET;ORAL	Prescription	No	AB
OXYMORPHONE HYDROCHLORIDE	OXYMORPHONE HYDROCHLORIDE	10MG	TABLET;ORAL	Prescription	No	AB

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

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