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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](#)
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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	AB
OXYMORPHONE HYDROCHLORIDE	OXYMORPHONE HYDROCHLORIDE	10MG		TABLET;ORAL	Prescription No	AB	AB

[Back to Top](#) | [Back to Previous Page](#) | [Back to Drugs@FDA Home](#)

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