Drug Details

Drug Name(s)	**OXYMORPHONE HYDROCHLORIDE**
FDA Application No.	**(ANDA) 203506**
Active Ingredient(s)	**OXYMORPHONE HYDROCHLORIDE**
Company	**SUN PHARM INDS LTD**
Original Approval or Tentative Approval Date	**April 24, 2015**

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

Products on Application (ANDA) #203506
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, EXTENDED RELEASE;ORAL	Prescription	No	AB
OXYMORPHONE HYDROCHLORIDE	OXYMORPHONE HYDROCHLORIDE	7.5MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB
OXYMORPHONE HYDROCHLORIDE	OXYMORPHONE HYDROCHLORIDE	10MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB
OXYMORPHONE HYDROCHLORIDE	OXYMORPHONE HYDROCHLORIDE	15MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB
OXYMORPHONE HYDROCHLORIDE	OXYMORPHONE HYDROCHLORIDE	20MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB
OXYMORPHONE HYDROCHLORIDE	OXYMORPHONE HYDROCHLORIDE	30MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB
OXYMORPHONE HYDROCHLORIDE	OXYMORPHONE HYDROCHLORIDE	40MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB

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