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## Drug Details

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

- [Therapeutic Equivalents](#)
- [Labels are not available](#)
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### Products on Application (ANDA) #203506

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

<a href="#">Drug Name</a>	<a href="#">Active Ingredients</a>	<a href="#">Strength</a>	<a href="#">Dosage Form/Route</a>	<a href="#">Marketing Status</a>	<a href="#">RLDTE Code</a>
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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