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## Drug Details

<b>Drug Name(s)</b>	<b>OPANA ER</b>
<b>FDA Application No.</b>	<b>(NDA) 021610</b>
<b>Active Ingredient(s)</b>	<b>OXYMORPHONE HYDROCHLORIDE</b>
<b>Company</b>	<b>ENDO PHARMS</b>
<b>Original Approval or Tentative Approval Date</b>	<b>June 22, 2006</b>
<b>Chemical Type</b>	<b>3 New dosage form</b>
<b>Review Classification</b>	<b>S Standard review drug</b>

- [There are no Therapeutic Equivalents](#)
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## Products on Application (NDA) #021610

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="#">RLD</a>	<a href="#">TE Code</a>
OPANA ER	OXYMORPHONE HYDROCHLORIDE	5MG	TABLET, EXTENDED RELEASE;ORAL	Discontinued	No	None
OPANA ER	OXYMORPHONE HYDROCHLORIDE	10MG	TABLET, EXTENDED RELEASE;ORAL	Discontinued	No	None
OPANA ER	OXYMORPHONE HYDROCHLORIDE	20MG	TABLET, EXTENDED RELEASE;ORAL	Discontinued	No	None
OPANA ER	OXYMORPHONE HYDROCHLORIDE	40MG	TABLET, EXTENDED RELEASE;ORAL	Discontinued	No	None
OPANA ER	OXYMORPHONE HYDROCHLORIDE	7.5MG	TABLET, EXTENDED RELEASE;ORAL	Discontinued	No	None
OPANA ER	OXYMORPHONE HYDROCHLORIDE	15MG	TABLET, EXTENDED RELEASE;ORAL	Discontinued	No	None
OPANA ER	OXYMORPHONE HYDROCHLORIDE	30MG	TABLET, EXTENDED RELEASE;ORAL	Discontinued	No	None

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