Drug Details

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

- **There are no Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **REMS[12]**
- **Label Information**
- **REMS[11]**

Products on Application (NDA) #021610
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
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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