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## Drug Details

<b>Drug Name(s)</b>	<b>OXYCODONE HYDROCHLORIDE</b>
<b>FDA Application No.</b>	<b>(ANDA) 091313</b>
<b>Active Ingredient(s)</b>	<b>OXYCODONE HYDROCHLORIDE</b>
<b>Company</b>	<b>COASTAL PHARMS</b>
<b>Original Approval or Tentative Approval Date</b>	<b>April 29, 2016</b>

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- **Labels are not available**

### Products on Application (ANDA) #091313

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<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>
OXYCODONE HYDROCHLORIDE	OXYCODONE HYDROCHLORIDE	5MG	TABLET;ORAL	Prescription	No	AB
OXYCODONE HYDROCHLORIDE	OXYCODONE HYDROCHLORIDE	15MG	TABLET;ORAL	Prescription	No	AB
OXYCODONE HYDROCHLORIDE	OXYCODONE HYDROCHLORIDE	30MG	TABLET;ORAL	Prescription	No	AB
OXYCODONE HYDROCHLORIDE	OXYCODONE HYDROCHLORIDE	10MG	TABLET;ORAL	Prescription	No	AB
OXYCODONE HYDROCHLORIDE	OXYCODONE HYDROCHLORIDE	20MG	TABLET;ORAL	Prescription	No	AB

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