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## Drug Details

<b>Drug Name(s)</b>	<b>TARGINIQ</b>
<b>FDA Application No.</b>	<b>(NDA) 205777</b>
<b>Active Ingredient(s)</b>	<b>NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE</b>
<b>Company</b>	<b>PURDUE PHARMA LP</b>
<b>Original Approval or Tentative Approval Date</b>	<b>July 23, 2014</b>
<b>Chemical Type</b>	<b>4 New combination</b>
<b>Review Classification</b>	<b>S Standard review drug</b>

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### Products on Application (NDA) #205777

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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="#">RLDTE Code</a>
TARGINIQ	NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE	5MG; 10MG	TABLET, EXTENDED RELEASE; ORAL	Discontinued	No None
TARGINIQ	NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE	10MG; 20MG	TABLET, EXTENDED RELEASE; ORAL	Discontinued	No None
TARGINIQ	NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE	20MG; 40MG	TABLET, EXTENDED RELEASE; ORAL	Discontinued	No None

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