Drugs@FDA
FDA Approved Drug Products

Drug Details

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

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

Products on Application (NDA) #205777
Click on a column header to re-sort the table:

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