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Drug Details

Drug Name(s)	MILNACIPRAN HYDROCHLORIDE
FDA Application No.	(ANDA) 205071
Active Ingredient(s)	MILNACIPRAN HYDROCHLORIDE
Company	LIBERTY PHARMA INC
Original Approval or Tentative Approval Date	January 27, 2016

- **There are no Therapeutic Equivalents**
- **Labels are not available**
- [Approval History, Letters, Reviews, and Related Documents](#)

Products on Application (ANDA) #205071

Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage	Form/Route	Marketing Status	RLD	TE Code
MILNACIPRAN HYDROCHLORIDE	MILNACIPRAN HYDROCHLORIDE	12.5MG		TABLET;ORAL	Discontinued	No	None
MILNACIPRAN HYDROCHLORIDE	MILNACIPRAN HYDROCHLORIDE	25MG		TABLET;ORAL	Discontinued	No	None
MILNACIPRAN HYDROCHLORIDE	MILNACIPRAN HYDROCHLORIDE	50MG		TABLET;ORAL	Discontinued	No	None
MILNACIPRAN HYDROCHLORIDE	MILNACIPRAN HYDROCHLORIDE	100MG		TABLET;ORAL	Discontinued	No	None

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