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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