Drug Details

Drug Name(s)	**MILNACIPRAN HYDROCHLORIDE**
FDA Application No.	**(ANDA) 205081**
Active Ingredient(s)	**MILNACIPRAN HYDROCHLORIDE**
Company	**AMNEAL PHARM**
Original Approval or Tentative Approval Date	**April 22, 2016**

- **There are no Therapeutic Equivalents**
- **Labels are not available**
- **Approval History, Letters, Reviews, and Related Documents**

Products on Application (ANDA) #205081
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	None (Tentative Approval)	No	None
MILNACIPRAN HYDROCHLORIDE	MILNACIPRAN HYDROCHLORIDE	25MG	TABLET;ORAL	None (Tentative Approval)	No	None
MILNACIPRAN HYDROCHLORIDE	MILNACIPRAN HYDROCHLORIDE	50MG	TABLET;ORAL	None (Tentative Approval)	No	None
MILNACIPRAN HYDROCHLORIDE	MILNACIPRAN HYDROCHLORIDE	100MG	TABLET;ORAL	None (Tentative Approval)	No	None

Back to Top | Back to Previous Page | Back to Drugs@FDA Home

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