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## Drug Details

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

- **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:

<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>
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

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