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## Drug Details

<b>Drug Name(s)</b>	<b>MEMANTINE HYDROCHLORIDE</b>
<b>FDA Application No.</b>	<b>(ANDA) 205825</b>
<b>Active Ingredient(s)</b>	<b>MEMANTINE HYDROCHLORIDE</b>
<b>Company</b>	<b>AMNEAL PHARMS</b>

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

### Products on Application (ANDA) #205825

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>
MEMANTINE HYDROCHLORIDE	MEMANTINE HYDROCHLORIDE	7MG	CAPSULE, EXTENDED RELEASE;ORAL	None (Tentative Approval)	No None
MEMANTINE HYDROCHLORIDE	MEMANTINE HYDROCHLORIDE	14MG	CAPSULE, EXTENDED RELEASE;ORAL	None (Tentative Approval)	No None
MEMANTINE HYDROCHLORIDE	MEMANTINE HYDROCHLORIDE	21MG	CAPSULE, EXTENDED RELEASE;ORAL	None (Tentative Approval)	No None
MEMANTINE HYDROCHLORIDE	MEMANTINE HYDROCHLORIDE	28MG	CAPSULE, EXTENDED RELEASE;ORAL	None (Tentative Approval)	No None

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