Drug Details

Drug Name(s)	**MEMANTINE HYDROCHLORIDE**
FDA Application No.	**(ANDA) 205825**
Active Ingredient(s)	**MEMANTINE HYDROCHLORIDE**
Company	**AMNEAL PHARMS**
Original Approval or Tentative Approval Date	**October 13, 2016**

- **Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Labels are not available**

Products on Application (ANDA) #205825
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
MEMANTINE HYDROCHLORIDE	MEMANTINE HYDROCHLORIDE	7MG	CAPSULE, EXTENDED RELEASE;ORAL	Prescription	No	AB
MEMANTINE HYDROCHLORIDE	MEMANTINE HYDROCHLORIDE	14MG	CAPSULE, EXTENDED RELEASE;ORAL	Prescription	No	AB
MEMANTINE HYDROCHLORIDE	MEMANTINE HYDROCHLORIDE	21MG	CAPSULE, EXTENDED RELEASE;ORAL	Prescription	No	AB
MEMANTINE HYDROCHLORIDE	MEMANTINE HYDROCHLORIDE	28MG	CAPSULE, EXTENDED RELEASE;ORAL	Prescription	No	AB

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