Drug Details

Drug Name(s)	**DESIPRAMINE HYDROCHLORIDE**
FDA Application No.	**(ANDA) 208105**
Active Ingredient(s)	**DESIPRAMINE HYDROCHLORIDE**
Company	**AMNEAL PHARMS CO**
Original Approval or Tentative Approval Date	**March 17, 2016**

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

Products on Application (ANDA) #208105
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
DESIPRAMINE HYDROCHLORIDE	DESIPRAMINE HYDROCHLORIDE	10MG	TABLET;ORAL	Prescription	No	AB
DESIPRAMINE HYDROCHLORIDE	DESIPRAMINE HYDROCHLORIDE	25MG	TABLET;ORAL	Prescription	No	AB
DESIPRAMINE HYDROCHLORIDE	DESIPRAMINE HYDROCHLORIDE	50MG	TABLET;ORAL	Prescription	No	AB
DESIPRAMINE HYDROCHLORIDE	DESIPRAMINE HYDROCHLORIDE	75MG	TABLET;ORAL	Prescription	No	AB
DESIPRAMINE HYDROCHLORIDE	DESIPRAMINE HYDROCHLORIDE	100MG	TABLET;ORAL	Prescription	No	AB
DESIPRAMINE HYDROCHLORIDE	DESIPRAMINE HYDROCHLORIDE	150MG	TABLET;ORAL	Prescription	No	AB

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