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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	RLDTE 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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