Drug Details

Drug Name(s)	**CLONIDINE HYDROCHLORIDE**
FDA Application No.	**(ANDA) 203320**
Active Ingredient(s)	**CLONIDINE HYDROCHLORIDE**
Company	**ACTAVIS ELIZABETH**
Original Approval or Tentative Approval Date	**June 15, 2015**

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

Products on Application (ANDA) #203320
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
CLONIDINE HYDROCHLORIDE	CLONIDINE HYDROCHLORIDE	0.1MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB1
CLONIDINE HYDROCHLORIDE	CLONIDINE HYDROCHLORIDE	0.2MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB1

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