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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

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- **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	RLDTE 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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