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Drug Details

Drug Name(s)	HYDRALAZINE HYDROCHLORIDE
FDA Application No.	(ANDA) 203110
Active Ingredient(s)	HYDRALAZINE HYDROCHLORIDE
Company	X-GEN PHARMS INC
Original Approval or Tentative Approval Date	June 29, 2015

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- **Labels are not available**

Products on Application (ANDA) #203110

Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength Dosage Form/Route	Marketing Status	RLDTE Code
HYDRALAZINE HYDROCHLORIDE	HYDRALAZINE HYDROCHLORIDE	20MG/ML INJECTABLE;INJECTION	Prescription	No AP

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