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Drug Details

Drug Name(s)	QUINAPRIL HYDROCHLORIDE
FDA Application No.	(ANDA) 205823
Active Ingredient(s)	QUINAPRIL HYDROCHLORIDE
Company	PRINSTON INC
Original Approval or Tentative Approval Date	September 15, 2016

- [Therapeutic Equivalents](#)
- [Approval History, Letters, Reviews, and Related Documents](#)
- **Labels are not available**

Products on Application (ANDA) #205823

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
QUINAPRIL HYDROCHLORIDE	QUINAPRIL HYDROCHLORIDE	EQ 5MG BASE	TABLET;ORAL	Prescription	No AB
QUINAPRIL HYDROCHLORIDE	QUINAPRIL HYDROCHLORIDE	EQ 10MG BASE	TABLET;ORAL	Prescription	No AB
QUINAPRIL HYDROCHLORIDE	QUINAPRIL HYDROCHLORIDE	EQ 20MG BASE	TABLET;ORAL	Prescription	No AB
QUINAPRIL HYDROCHLORIDE	QUINAPRIL HYDROCHLORIDE	EQ 40MG BASE	TABLET;ORAL	Prescription	No AB

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