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	RLD	TE 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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