Drug Details

Drug Name(s)	**SOTALOL HYDROCHLORIDE**
FDA Application No.	**(ANDA) 207428**
Active Ingredient(s)	**SOTALOL HYDROCHLORIDE**
Company	**BEXIMCO PHARMS USA**
Original Approval or Tentative Approval Date	**October 21, 2016**

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

Products on Application (ANDA) #207428
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
SOTALOL HYDROCHLORIDE	SOTALOL HYDROCHLORIDE	80MG	TABLET;ORAL	Prescription	No	AB1
SOTALOL HYDROCHLORIDE	SOTALOL HYDROCHLORIDE	120MG	TABLET;ORAL	Prescription	No	AB1
SOTALOL HYDROCHLORIDE	SOTALOL HYDROCHLORIDE	160MG	TABLET;ORAL	Prescription	No	AB1

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