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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

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- **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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