Drug Details

Drug Name(s)	**NEBIVOLOL HYDROCHLORIDE**
FDA Application No.	**(ANDA) 203683**
Active Ingredient(s)	**NEBIVOLOL HYDROCHLORIDE**
Company	**WATSON LABS INC**
Original Approval or Tentative Approval Date	**November 27, 2015**

- **There are no Therapeutic Equivalents**
- **Labels are not available**
- **Approval History, Letters, Reviews, and Related Documents**

Products on Application (ANDA) #203683
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
NEBIVOLOL HYDROCHLORIDE	NEBIVOLOL HYDROCHLORIDE	EQ 2.5MG BASE	TABLET;ORAL	Discontinued	No	None
NEBIVOLOL HYDROCHLORIDE	NEBIVOLOL HYDROCHLORIDE	EQ 5MG BASE	TABLET;ORAL	Discontinued	No	None
NEBIVOLOL HYDROCHLORIDE	NEBIVOLOL HYDROCHLORIDE	EQ 10MG BASE	TABLET;ORAL	Discontinued	No	None
NEBIVOLOL HYDROCHLORIDE	NEBIVOLOL HYDROCHLORIDE	EQ 20MG BASE	TABLET;ORAL	Discontinued	No	None

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