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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	RLDTE 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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