Drugs@FDA
FDA Approved Drug Products

Drug Details

Drug Name(s)	**NADOLOL**
FDA Application No.	**(ANDA) 201893**
Active Ingredient(s)	**NADOLOL**
Company	**ORION CORP ORION**
Original Approval or Tentative Approval Date	**September 16, 2015**

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

Products on Application (ANDA) #201893
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
NADOLOL	NADOLOL	40MG	TABLET;ORAL	Prescription	No	AB
NADOLOL	NADOLOL	80MG	TABLET;ORAL	Prescription	No	AB

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