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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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