Drugs@FDA
FDA Approved Drug Products



## Drug Details

| | |
|---|---|
| **Drug Name(s)** | **NITROFURANTOIN** |
| **FDA Application No.** | **(ANDA) 201722** |
| **Active Ingredient(s)** | **NITROFURANTOIN, MACROCRYSTALLINE** |
| **Company** | **SUN PHARM INDS** |
| **Original Approval or Tentative Approval Date** | **February 16, 2016** |

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

**Products on Application (ANDA) #201722**
**Click on a column header to re-sort the table:**

| Drug Name | Active Ingredients | Strength | Dosage Form/Route | Marketing Status | RLD | TE Code |
|---|---|---|---|---|---|---|
| NITROFURANTOIN | NITROFURANTOIN, MACROCRYSTALLINE | 25MG | CAPSULE;ORAL | Prescription | No | AB |
| NITROFURANTOIN | NITROFURANTOIN, MACROCRYSTALLINE | 50MG | CAPSULE;ORAL | Prescription | No | AB |
| NITROFURANTOIN | NITROFURANTOIN, MACROCRYSTALLINE | 100MG | CAPSULE;ORAL | Prescription | No | AB |

Back to Top | Back to Previous Page | Back to Drugs@FDA Home



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