## Drug Details

| | |
|---|---|
| **Drug Name(s)** | **IRINOTECAN HYDROCHLORIDE** |
| **FDA Application No.** | **(ANDA) 203380** |
| **Active Ingredient(s)** | **IRINOTECAN HYDROCHLORIDE** |
| **Company** | **QILU PHARM CO LTD** |
| **Original Approval or Tentative Approval Date** | **May 3, 2016** |

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

**Products on Application (ANDA) #203380**
**Click on a column header to re-sort the table:**

| Drug Name | Active Ingredients | Strength | Dosage Form/Route | Marketing Status | RLD | TE Code |
|---|---|---|---|---|---|---|
| IRINOTECAN HYDROCHLORIDE | IRINOTECAN HYDROCHLORIDE | 40MG/2ML (20MG/ML) | INJECTABLE;INJECTION | Prescription | No | AP |
| IRINOTECAN HYDROCHLORIDE | IRINOTECAN HYDROCHLORIDE | 100MG/5ML (20MG/ML) | INJECTABLE;INJECTION | Prescription | No | AP |
| IRINOTECAN HYDROCHLORIDE | IRINOTECAN HYDROCHLORIDE | 300MG/15ML (20MG/ML) | INJECTABLE;INJECTION | Prescription | No | AP |

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