Drugs@FDA
FDA Approved Drug Products



## Drug Details

| | |
|---|---|
| **Drug Name(s)** | **DULOXETINE HYDROCHLORIDE** |
| **FDA Application No.** | **(ANDA) 204343** |
| **Active Ingredient(s)** | **DULOXETINE HYDROCHLORIDE** |
| **Company** | **HETERO LABS LTD III** |
| **Original Approval or Tentative Approval Date** | **August 4, 2016** |

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

**Products on Application (ANDA) #204343**
**Click on a column header to re-sort the table:**

| Drug Name | Active Ingredients | Strength | Dosage Form/Route | Marketing Status | RLD | TE Code |
|---|---|---|---|---|---|---|
| DULOXETINE HYDROCHLORIDE | DULOXETINE HYDROCHLORIDE | EQ 20MG BASE | CAPSULE, DELAYED REL PELLETS;ORAL | Prescription | No | AB |
| DULOXETINE HYDROCHLORIDE | DULOXETINE HYDROCHLORIDE | EQ 30MG BASE | CAPSULE, DELAYED REL PELLETS;ORAL | Prescription | No | AB |
| DULOXETINE HYDROCHLORIDE | DULOXETINE HYDROCHLORIDE | EQ 60MG BASE | CAPSULE, DELAYED REL PELLETS;ORAL | Prescription | No | AB |

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