## Drug Details

| | |
|---|---|
| **Drug Name(s)** | **ACZONE** |
| **FDA Application No.** | **(NDA) 207154** |
| **Active Ingredient(s)** | **DAPSONE** |
| **Company** | **ALLERGAN** |
| **Original Approval or Tentative Approval Date** | **February 24, 2016** |
| **Chemical Type** | **5 New formulation or new manufacturer** |

- **There are no Therapeutic Equivalents**
- **Labels are not available**
- **Approval History, Letters, Reviews, and Related Documents**

**Products on Application (NDA) #207154**
**Click on a column header to re-sort the table:**

| Drug Name | Active Ingredients | Strength | Dosage Form/Route | Marketing Status | RLD | TE Code |
|---|---|---|---|---|---|---|
| ACZONE | DAPSONE | 7.5% | GEL; TOPICAL | Prescription | TBD [11] | None |

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