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Drug Details

| | |
|---|--------------------------------|
| Drug Name(s) | BUPROPION HYDROCHLORIDE |
| FDA Application No. | (ANDA) 206122 |
| Active Ingredient(s) | BUPROPION HYDROCHLORIDE |
| Company | SCIEGEN PHARMS INC |
| Original Approval or Tentative Approval Date | August 17, 2016 |

- [Therapeutic Equivalents](#)
- [Approval History, Letters, Reviews, and Related Documents](#)
- **Labels are not available**

Products on Application (ANDA) #206122

Click on a column header to re-sort the table:

| Drug Name | Active Ingredients | Strength | Dosage Form/Route | Marketing Status | RLDTE Code |
|---------------------------|------------------------------------|--------------------------|-----------------------------------|----------------------------------|----------------------------|
| BUPROPION HYDROCHLORIDE | BUPROPION HYDROCHLORIDE | 150MG | TABLET, EXTENDED RELEASE; ORAL | Prescription No | AB2 |

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