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	RLD	TE Code
BUPROPION HYDROCHLORIDE	BUPROPION HYDROCHLORIDE	150MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB2

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

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