Drug Details

Drug Name(s)	**BUDESONIDE**
FDA Application No.	**(ANDA) 206623**
Active Ingredient(s)	**BUDESONIDE**
Company	**ACTAVIS ELIZABETH**
Original Approval or Tentative Approval Date	**April 8, 2016**

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

Products on Application (ANDA) #206623
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
BUDESONIDE	BUDESONIDE	3MG	CAPSULE;ORAL	Prescription	No	AB

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