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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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