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Drug Details

Drug Name(s)	DESOXIMETASONE
FDA Application No.	(ANDA) 205082
Active Ingredient(s)	DESOXIMETASONE
Company	ACTAVIS MID ATLANTIC
Original Approval or Tentative Approval Date	September 4, 2015

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

Products on Application (ANDA) #205082

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
DESOXIMETASONE	DESOXIMETASONE	0.25%	CREAM;TOPICAL	Prescription	No	AB

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