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Drug Details

Drug Name(s)	DESOXIMETASONE
FDA Application No.	(ANDA) 204675
Active Ingredient(s)	DESOXIMETASONE
Company	GROUPE PARIMA INC
Original Approval or Tentative Approval Date	August 15, 2016

- [Therapeutic Equivalents](#)
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- **Labels are not available**

Products on Application (ANDA) #204675

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	0.05%	GEL;TOPICAL	Prescription	No	AB

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