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Drug Details

Drug Name(s)	MOMETASONE FUROATE
FDA Application No.	(ANDA) 091161
Active Ingredient(s)	MOMETASONE FUROATE
Company	APOTEX INC
Original Approval or Tentative Approval Date	March 22, 2016

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

Products on Application (ANDA) #091161

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
MOMETASONE FUROATE	MOMETASONE FUROATE	EQ 0.05MG BASE/SPRAY	SPRAY, METERED;NASAL	Prescription	No AB

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