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Drug Details

Drug Name(s)	CALCIUM ACETATE
FDA Application No.	(ANDA) 203179
Active Ingredient(s)	CALCIUM ACETATE
Company	NOSTRUM LABS INC
Original Approval or Tentative Approval Date	October 26, 2015

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

Products on Application (ANDA) #203179

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD TE Code
CALCIUM ACETATE	CALCIUM ACETATE	EQ 169MG CALCIUM	CAPSULE;ORAL	Prescription	No AB

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