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Drug Details

Drug Name(s)	FAMOTIDINE
FDA Application No.	(ANDA) 206530
Active Ingredient(s)	FAMOTIDINE
Company	AUROBINDO PHARMA LTD
Original Approval or Tentative Approval Date	December 22, 2015

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

Products on Application (ANDA) #206530

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
FAMOTIDINE	FAMOTIDINE	20MG	TABLET;ORAL	Prescription	No	AB
FAMOTIDINE	FAMOTIDINE	40MG	TABLET;ORAL	Prescription	No	AB

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