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Drug Details

Drug Name(s)	ETOMIDATE
FDA Application No.	(ANDA) 202354
Active Ingredient(s)	ETOMIDATE
Company	HIKMA FARMACEUTICA
Original Approval or Tentative Approval Date	February 25, 2016

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- **Labels are not available**

Products on Application (ANDA) #202354

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
ETOMIDATE	ETOMIDATE	2MG/ML	INJECTABLE;INJECTION	Prescription	No	AP

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