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Drug Details

Drug Name(s)	DONEPEZIL HYDROCHLORIDE
FDA Application No.	(ANDA) 203419
Active Ingredient(s)	DONEPEZIL HYDROCHLORIDE
Company	INDICUS PHARMA
Original Approval or Tentative Approval Date	April 12, 2016

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- **Labels are not available**

Products on Application (ANDA) #203419

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
DONEPEZIL HYDROCHLORIDE	DONEPEZIL HYDROCHLORIDE	23MG	TABLET;ORAL	Prescription	No	AB

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