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Drug Details

Drug Name(s)	DONEPEZIL HYDROCHLORIDE
FDA Application No.	(ANDA) 204293
Active Ingredient(s)	DONEPEZIL HYDROCHLORIDE
Company	SUN PHARM INDS LTD
Original Approval or Tentative Approval Date	June 5, 2015

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

Products on Application (ANDA) #204293

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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