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Drug Details

Drug Name(s)	GALANTAMINE HYDROBROMIDE
FDA Application No.	(ANDA) 204895
Active Ingredient(s)	GALANTAMINE HYDROBROMIDE
Company	AUROBINDO PHARMA LTD
Original Approval or Tentative Approval Date	August 5, 2016

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

Products on Application (ANDA) #204895

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
GALANTAMINE HYDROBROMIDE	GALANTAMINE HYDROBROMIDE	EQ 8MG BASE	CAPSULE, EXTENDED RELEASE;ORAL	Prescription No	AB
GALANTAMINE HYDROBROMIDE	GALANTAMINE HYDROBROMIDE	EQ 16MG BASE	CAPSULE, EXTENDED RELEASE;ORAL	Prescription No	AB
GALANTAMINE HYDROBROMIDE	GALANTAMINE HYDROBROMIDE	EQ 24MG BASE	CAPSULE, EXTENDED RELEASE;ORAL	Prescription No	AB

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