Drugs@FDA
FDA Approved Drug Products

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	RLD	TE 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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