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Drug Details

Drug Name(s)	RALOXIFENE HYDROCHLORIDE
FDA Application No.	(ANDA) 204491
Active Ingredient(s)	RALOXIFENE HYDROCHLORIDE
Company	GLENMARK PHARMS LTD
Original Approval or Tentative Approval Date	March 22, 2016

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

Products on Application (ANDA) #204491

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
RALOXIFENE HYDROCHLORIDE	RALOXIFENE HYDROCHLORIDE	60MG	TABLET;ORAL	Prescription	No	AB

[Back to Top](#) | [Back to Previous Page](#) | [Back to Drugs@FDA Home](#)

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