Drug Details

Drug Name(s)	**RISEDRONATE SODIUM**
FDA Application No.	**(ANDA) 203533**
Active Ingredient(s)	**RISEDRONATE SODIUM**
Company	**MACLEODS PHARMS LTD**
Original Approval or Tentative Approval Date	**December 9, 2015**

- **Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Labels are not available**

Products on Application (ANDA) #203533
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
RISEDRONATE SODIUM	RISEDRONATE SODIUM	5MG	TABLET;ORAL	Prescription	No	AB
RISEDRONATE SODIUM	RISEDRONATE SODIUM	30MG	TABLET;ORAL	Prescription	No	AB
RISEDRONATE SODIUM	RISEDRONATE SODIUM	35MG	TABLET;ORAL	None (Tentative Approval)	No	None

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