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## Drug Details

<b>Drug Name(s)</b>	QUININE SULFATE
<b>FDA Application No.</b>	(ANDA) 203112
<b>Active Ingredient(s)</b>	QUININE SULFATE
<b>Company</b>	LUPIN LTD
<b>Original Approval or Tentative Approval Date</b>	April 24, 2015

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

### Products on Application (ANDA) #203112

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
QUININE SULFATE	QUININE SULFATE	324MG	CAPSULE;ORAL	Prescription	No	AB

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