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Drug Details

Drug Name(s)	LITHIUM CARBONATE
FDA Application No.	(ANDA) 204445
Active Ingredient(s)	LITHIUM CARBONATE
Company	ALEMBIC PHARMS LTD
Original Approval or Tentative Approval Date	June 10, 2015

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

Products on Application (ANDA) #204445

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD TE Code
LITHIUM CARBONATE	LITHIUM CARBONATE	300MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No AB

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