Drug Details

Drug Name(s)	**PARICALCITOL**
FDA Application No.	**(ANDA) 206710**
Active Ingredient(s)	**PARICALCITOL**
Company	**LOTUS PHARM CO LTD**
Original Approval or Tentative Approval Date	**February 24, 2016**

- **There are no Therapeutic Equivalents**
- **Labels are not available**
- **Approval History, Letters, Reviews, and Related Documents**

Products on Application (ANDA) #206710
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
PARICALCITOL	PARICALCITOL	1MCG	CAPSULE;ORAL	None (Tentative Approval)	No	None
PARICALCITOL	PARICALCITOL	2MCG	CAPSULE;ORAL	None (Tentative Approval)	No	None
PARICALCITOL	PARICALCITOL	4MCG	CAPSULE;ORAL	None (Tentative Approval)	No	None

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