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Drug Details

Drug Name(s)	LANSOPRAZOLE
FDA Application No.	(ANDA) 203203
Active Ingredient(s)	LANSOPRAZOLE
Company	ANCHEN PHARMS
Original Approval or Tentative Approval Date	July 26, 2016

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

Products on Application (ANDA) #203203

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
LANSOPRAZOLE	LANSOPRAZOLE	15MG	CAPSULE, DELAYED REL PELLETS;ORAL	Prescription	No	AB
LANSOPRAZOLE	LANSOPRAZOLE	30MG	CAPSULE, DELAYED REL PELLETS;ORAL	Prescription	No	AB

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