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Drug Details

Drug Name(s)	OMEPRAZOLE AND SODIUM BICARBONATE
FDA Application No.	(ANDA) 204922
Active Ingredient(s)	OMEPRAZOLE; SODIUM BICARBONATE
Company	AUROLIFE PHARMA LLC
Original Approval or Tentative Approval Date	August 21, 2016

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

Products on Application (ANDA) #204922

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
OMEPRAZOLE AND SODIUM BICARBONATE	OMEPRAZOLE; SODIUM BICARBONATE	20MG; 1.1GM	CAPSULE;ORAL	Prescription No	AB
OMEPRAZOLE AND SODIUM BICARBONATE	OMEPRAZOLE; SODIUM BICARBONATE	40MG; 1.1GM	CAPSULE;ORAL	Prescription No	AB

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