Drugs@FDA
FDA Approved Drug Products

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**

- **Therapeutic Equivalents**
- **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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