Drug Details

Drug Name(s)	**LOMAIRA**
FDA Application No.	**(ANDA) 203495**
Active Ingredient(s)	**PHENTERMINE HYDROCHLORIDE**
Company	**AVANTHI INC**
Original Approval or Tentative Approval Date	**September 13, 2016**

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

Products on Application (ANDA) #203495
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
LOMAIRA	PHENTERMINE HYDROCHLORIDE	8MG	TABLET;ORAL	Prescription	Yes	None

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