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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

[Back to Top](#) | [Back to Previous Page](#) | [Back to Drugs@FDA Home](#)

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