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Drug Details

Drug Name(s)	METHAMPHETAMINE HYDROCHLORIDE
FDA Application No.	(ANDA) 203846
Active Ingredient(s)	METHAMPHETAMINE HYDROCHLORIDE
Company	ROXANE
Original Approval or Tentative Approval Date	November 17, 2015

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

Products on Application (ANDA) #203846

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

Drug Name	Active Ingredients	Strength Dosage Form/Route	Marketing Status	RLDTE Code
METHAMPHETAMINE HYDROCHLORIDE	METHAMPHETAMINE HYDROCHLORIDE	5MG TABLET;ORAL	Prescription No	AA

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U.S. Food and Drug Administration
10903 New Hampshire Avenue
Silver Spring, MD 20993
Ph. 1-888-INFO-FDA (1-888-463-6332)

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