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Drug Details

Drug Name(s)	DEXTROAMPHETAMINE SULFATE
FDA Application No.	(ANDA) 203548
Active Ingredient(s)	DEXTROAMPHETAMINE SULFATE
Company	AVANTHI INC
Original Approval or Tentative Approval Date	November 24, 2015

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

Products on Application (ANDA) #203548

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

Drug Name	Active Ingredients	Strength	Dosage	Form/Route	Marketing Status	RLDTE Code
DEXTROAMPHETAMINE SULFATE	DEXTROAMPHETAMINE SULFATE	5MG		TABLET;ORAL	Prescription	No AA
DEXTROAMPHETAMINE SULFATE	DEXTROAMPHETAMINE SULFATE	10MG		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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