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Drug Details

Drug Name(s)	AMMONIA N 13
FDA Application No.	(ANDA) 204515
Active Ingredient(s)	AMMONIA N-13
Company	NCM USA BRONX LLC
Original Approval or Tentative Approval Date	February 4, 2015

- **There are no Therapeutic Equivalents**
- **Labels are not available**
- [Approval History, Letters, Reviews, and Related Documents](#)

Products on Application (ANDA) #204515

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD TE Code
AMMONIA N 13	AMMONIA N-13	3.75-260mCi/mL	INJECTABLE;INTRAVENOUS	Prescription	No None

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