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Drug Details

Drug Name(s)	LEVOTHYROXINE SODIUM
FDA Application No.	(ANDA) 206163
Active Ingredient(s)	LEVOTHYROXINE SODIUM
Company	FERA PHARMS LLC
Original Approval or Tentative Approval Date	June 29, 2016

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

Products on Application (ANDA) #206163

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
LEVOTHYROXINE SODIUM	LEVOTHYROXINE SODIUM	100MCG/VIAL	POWDER;INTRAVENOUS	Prescription	No	AP
LEVOTHYROXINE SODIUM	LEVOTHYROXINE SODIUM	500MCG/VIAL	POWDER;INTRAVENOUS	Prescription	No	AP

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