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Drug Details

Drug Name(s)	ZOLEDRONIC ACID
FDA Application No.	(ANDA) 090621
Active Ingredient(s)	ZOLEDRONIC ACID
Company	HOSPIRA INC
Original Approval or Tentative Approval Date	March 19, 2015

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- **Labels are not available**

Products on Application (ANDA) #090621

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD TE Code
ZOLEDRONIC ACID	ZOLEDRONIC ACID	EQ 4MG BASE/5ML	INJECTABLE;IV (INFUSION)	Prescription	No AP

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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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