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Drug Details

Drug Name(s)	SEVOFLURANE
FDA Application No.	(ANDA) 203793
Active Ingredient(s)	SEVOFLURANE
Company	SHANGHAI HENGRUI
Original Approval or Tentative Approval Date	November 3, 2015

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- **Labels are not available**

Products on Application (ANDA) #203793

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
SEVOFLURANE	SEVOFLURANE	100%	LIQUID;INHALATION	Prescription	No	AN

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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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