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Drug Details

Drug Name(s)	LEVOCETIRIZINE DIHYDROCHLORIDE
FDA Application No.	(ANDA) 203027
Active Ingredient(s)	LEVOCETIRIZINE DIHYDROCHLORIDE
Company	APOTEX INC
Original Approval or Tentative Approval Date	February 13, 2015

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Products on Application (ANDA) #203027
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
LEVOCETIRIZINE DIHYDROCHLORIDE	LEVOCETIRIZINE DIHYDROCHLORIDE	5MG	TABLET;ORAL	Prescription No	AB

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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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