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Drug Details

Drug Name(s)	CETIRIZINE HYDROCHLORIDE ALLERGY
FDA Application No.	(ANDA) 207235
Active Ingredient(s)	CETIRIZINE HYDROCHLORIDE
Company	APOTEX INC
Original Approval or Tentative Approval Date	August 15, 2016

- [Other OTC Drugs with the same Active Ingredient, Strength and Dosage Form/Route](#)
- [Approval History, Letters, Reviews, and Related Documents](#)
- Labels are not available

Products on Application (ANDA) #207235
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
CETIRIZINE HYDROCHLORIDE ALLERGY	CETIRIZINE HYDROCHLORIDE	10MG	CAPSULE;ORAL	Over-the-counter	No None

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