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Drug Details

Drug Name(s)	CYCLOPENTOLATE HYDROCHLORIDE
FDA Application No.	(ANDA) 205937
Active Ingredient(s)	CYCLOPENTOLATE HYDROCHLORIDE
Company	AKORN INC
Original Approval or Tentative Approval Date	December 9, 2015

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- **Labels are not available**

Products on Application (ANDA) #205937

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
CYCLOPENTOLATE HYDROCHLORIDE	CYCLOPENTOLATE HYDROCHLORIDE	0.5%	SOLUTION/DROPS;OPHTHALMIC	Prescription No	AT

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