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Drug Details

Drug Name(s)	DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE
FDA Application No.	(ANDA) 202975
Active Ingredient(s)	DUTASTERIDE; TAMSULOSIN HYDROCHLORIDE
Company	ACTAVIS LABS FL INC
Original Approval or Tentative Approval Date	November 20, 2015

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Products on Application (ANDA) #202975

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE	DUTASTERIDE; TAMSULOSIN HYDROCHLORIDE	0.5MG; 0.4MG	CAPSULE;ORAL	Prescription No	AB

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