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Drug Details

Drug Name(s)	DUTASTERIDE
FDA Application No.	(ANDA) 202421
Active Ingredient(s)	DUTASTERIDE
Company	VINTAGE PHARMS LLC
Original Approval or Tentative Approval Date	November 20, 2015

- [Therapeutic Equivalents](#)
- [Approval History, Letters, Reviews, and Related Documents](#)
- Labels are not available

Products on Application (ANDA) #202421

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
DUTASTERIDE	DUTASTERIDE	0.5MG	CAPSULE;ORAL	Prescription	No	AB

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