Drug Details

Drug Name(s)	**TELMISARTAN AND HYDROCHLOROTHIAZIDE**
FDA Application No.	**(ANDA) 204169**
Active Ingredient(s)	**HYDROCHLOROTHIAZIDE; TELMISARTAN**
Company	**MACLEODS PHARMS LTD**
Original Approval or Tentative Approval Date	**November 2, 2015**

- **Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Labels are not available**

Products on Application (ANDA) #204169
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
TELMISARTAN AND HYDROCHLOROTHIAZIDE	HYDROCHLOROTHIAZIDE; TELMISARTAN	12.5MG; 40MG	TABLET;ORAL	Prescription	No AB
TELMISARTAN AND HYDROCHLOROTHIAZIDE	HYDROCHLOROTHIAZIDE; TELMISARTAN	12.5MG; 80MG	TABLET;ORAL	Prescription	No AB
TELMISARTAN AND HYDROCHLOROTHIAZIDE	HYDROCHLOROTHIAZIDE; TELMISARTAN	25MG; 80MG	TABLET;ORAL	Prescription	No AB

Back to Top | Back to Previous Page | Back to Drugs@FDA Home

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