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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

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