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Drug Details

Drug Name(s)	OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE
FDA Application No.	(ANDA) 203580
Active Ingredient(s)	AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL
Company	TORRENT PHARMS LTD
Original Approval or Tentative Approval Date	October 26, 2016

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

Products on Application (ANDA) #203580
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE	AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE;	EQ 5MG BASE; 12.5MG; 20MG	TABLET;ORAL	Prescription No	AB
OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE	OLMESARTAN MEDOXOMIL AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE;	EQ 5MG BASE; 12.5MG; 40MG	TABLET;ORAL	Prescription No	AB
OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE	OLMESARTAN MEDOXOMIL AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE;	EQ 5MG BASE; 25MG; 40MG	TABLET;ORAL	Prescription No	AB
OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE	OLMESARTAN MEDOXOMIL AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE;	EQ 10MG BASE; 12.5MG; 40MG	TABLET;ORAL	Prescription No	AB
OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE	OLMESARTAN MEDOXOMIL AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE;	EQ 10MG BASE; 25MG; 40MG	TABLET;ORAL	Prescription No	AB

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