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	RLD	TE Code
OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE	AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL	EQ 5MG BASE; 12.5MG; 20MG	TABLET;ORAL	Prescription	No	AB
OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE	AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL	EQ 5MG BASE; 12.5MG; 40MG	TABLET;ORAL	Prescription	No	AB
OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE	AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL	EQ 5MG BASE; 25MG; 40MG	TABLET;ORAL	Prescription	No	AB
OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE	AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL	EQ 10MG BASE; 12.5MG; 40MG	TABLET;ORAL	Prescription	No	AB
OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE	AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL	EQ 10MG BASE; 25MG; 40MG	TABLET;ORAL	Prescription	No	AB

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