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Drug Details

Drug Name(s)	AMLODIPINE AND OLMESARTAN MEDOXOMIL
FDA Application No.	(ANDA) 091154
Active Ingredient(s)	AMLODIPINE BESYLATE; OLMESARTAN MEDOXOMIL
Company	TEVA PHARMS USA
Original Approval or Tentative Approval Date	October 26, 2016

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Products on Application (ANDA) #091154

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

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

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