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Drug Details

Drug Name(s)	AMLODIPINE BESYLATE
FDA Application No.	(ANDA) 207821
Active Ingredient(s)	AMLODIPINE BESYLATE
Company	POLYGEN PHARMS
Original Approval or Tentative Approval Date	July 11, 2016

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

Products on Application (ANDA) #207821

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
AMLODIPINE BESYLATE	AMLODIPINE BESYLATE	EQ 2.5MG BASE	TABLET;ORAL	Prescription	No	AB
AMLODIPINE BESYLATE	AMLODIPINE BESYLATE	EQ 5MG BASE	TABLET;ORAL	Prescription	No	AB
AMLODIPINE BESYLATE	AMLODIPINE BESYLATE	EQ 10MG BASE	TABLET;ORAL	Prescription	No	AB

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