Drug Details

Drug Name(s)	**ARGATROBAN IN 0.9% SODIUM CHLORIDE**
FDA Application No.	**(NDA) 206769**
Active Ingredient(s)	**ARGATROBAN**
Company	**TEVA PHARMS USA**
Original Approval or Tentative Approval Date	**December 15, 2014**
Chemical Type	**5 New formulation or new manufacturer**
Review Classification	**S Standard review drug**

- **There are no Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Label Information**

Products on Application (NDA) #206769
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
ARGATROBAN IN 0.9% SODIUM CHLORIDE	ARGATROBAN	250MG/250ML (1MG/ML)	INJECTABLE;IV (INFUSION)	Prescription	No	None

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