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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](#)
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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	RLDTE Code
ARGATROBAN IN 0.9% SODIUM CHLORIDE	ARGATROBAN	250MG/250ML (1MG/ML)	INJECTABLE;IV (INFUSION)	Prescription No	None

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