Drug Details

Drug Name(s)	**SODIUM PHENYLACETATE AND SODIUM BENZOATE**
FDA Application No.	**(ANDA) 205880**
Active Ingredient(s)	**SODIUM BENZOATE; SODIUM PHENYLACETATE**
Company	**NAVINTA LLC**
Original Approval or Tentative Approval Date	**August 4, 2016**

- **There are no Therapeutic Equivalents**
- **Labels are not available**
- **Approval History, Letters, Reviews, and Related Documents**

Products on Application (ANDA) #205880
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
SODIUM PHENYLACETATE AND SODIUM BENZOATE	SODIUM BENZOATE; SODIUM PHENYLACETATE	10%; 10% (5GM/50ML; 5GM/50ML)	SOLUTION;IV (INFUSION)	Prescription	No	AP

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