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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	RLDTE Code
SODIUM PHENYLACETATE AND SODIUM BENZOATE	SODIUM BENZOATE; SODIUM PHENYLACETATE	10%; 10% (5GM/50ML; 5GM/50ML)	SOLUTION;IV (INFUSION)	Prescription No	AP

[Back to Top](#) | [Back to Previous Page](#) | [Back to Drugs@FDA Home](#)

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