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Drug Details

| | |
|---|-------------------------------|
| Drug Name(s) | SODIUM PHENYL BUTYRATE |
| FDA Application No. | (ANDA) 204395 |
| Active Ingredient(s) | SODIUM PHENYL BUTYRATE |
| Company | PAR PHARM |
| Original Approval or Tentative Approval Date | May 6, 2016 |

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

Products on Application (ANDA) #204395

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

| Drug Name | Active Ingredients | Strength | Dosage Form/Route | Marketing Status | RLD TE Code |
|---------------------------|------------------------------------|--------------------------|-----------------------------------|----------------------------------|-----------------------------|
| SODIUM PHENYL BUTYRATE | SODIUM PHENYL BUTYRATE | 500MG | TABLET;ORAL | Prescription | No AB |

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

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