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## Drug Details

|                                                     |                                                 |
|-----------------------------------------------------|-------------------------------------------------|
| <b>Drug Name(s)</b>                                 | <b>SODIUM PHENYLACETATE AND SODIUM BENZOATE</b> |
| <b>FDA Application No.</b>                          | <b>(ANDA) 207096</b>                            |
| <b>Active Ingredient(s)</b>                         | <b>SODIUM BENZOATE; SODIUM PHENYLACETATE</b>    |
| <b>Company</b>                                      | <b>AILEX PHARMS PVT LTD</b>                     |
| <b>Original Approval or Tentative Approval Date</b> | <b>February 24, 2016</b>                        |

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

## Products on Application (ANDA) #207096

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         |

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