



**Start Over**

### Drug Details

|   |                                     |
|---|-------------------------------------|
| <b>Drug Name(s)</b>                                 | <b>OCUFEN</b>                       |
| <b>FDA Application No.</b>                          | <b>(NDA) 019404</b>                 |
| <b>Active Ingredient(s)</b>                         | <b>FLURBIPROFEN SODIUM</b>          |
| <b>Company</b>                                      | <b>ALLERGAN</b>                     |
| <b>Original Approval or Tentative Approval Date</b> | <b>December 31, 1986</b>            |
| <b>Chemical Type</b>                                | <b>1 New molecular entity (NME)</b> |
| <b>Review Classification</b>                        | <b>P Priority review drug</b>       |

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**Products on Application (NDA) #019404**  
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|---------------------------|------------------------------------|--------------------------|-----------------------------------|----------------------------------|-----------------------------|
| OCUFEN                    | FLURBIPROFEN SODIUM                | 0.03%                    | SOLUTION/DROPS;OPHTHALMIC         | Prescription                     | Yes AT                      |

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