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## 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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- [Approval History, Letters, Reviews, and Related Documents](#)

## Products on Application (NDA) #019404

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OCUFEN	FLURBIPROFEN SODIUM	0.03%	SOLUTION/DROPS;OPHTHALMIC	Prescription	Yes	AT

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