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[Email Link](#)

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[Back to Search Results](#)

Drug Details

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

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

Products on Application (NDA) #019404

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
OCUFEN	FLURBIPROFEN SODIUM	0.03%	SOLUTION/DROPS;OPHTHALMIC	Prescription	Yes	AT

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

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