U.S. Department of Health and Human Services

Drugs@FDA
FDA Approved Drug Products

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**

- **Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Label Information**

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

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