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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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Products on Application (NDA) #019404
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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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U.S. Food and Drug Administration
10903 New Hampshire Avenue
Silver Spring, MD 20993
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