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Drug Details

Drug Name(s)	MOXIFLOXACIN HYDROCHLORIDE
FDA Application No.	(ANDA) 202525
Active Ingredient(s)	MOXIFLOXACIN HYDROCHLORIDE
Company	WATSON LABS INC
Original Approval or Tentative Approval Date	March 6, 2015

- [Therapeutic Equivalents](#)
- [Approval History, Letters, Reviews, and Related Documents](#)
- **Labels are not available**

Products on Application (ANDA) #202525

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RDTE Code
MOXIFLOXACIN HYDROCHLORIDE	MOXIFLOXACIN HYDROCHLORIDE	EQ 0.5% BASE	SOLUTION/DROPS;OPHTHALMIC	Prescription No	AT

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