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Drug Details

Drug Name(s)	HYDROMORPHONE HYDROCHLORIDE
FDA Application No.	(ANDA) 205814
Active Ingredient(s)	HYDROMORPHONE HYDROCHLORIDE
Company	AUROLIFE PHARMA LLC
Original Approval or Tentative Approval Date	May 13, 2016

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

Products on Application (ANDA) #205814

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
HYDROMORPHONE HYDROCHLORIDE	HYDROMORPHONE HYDROCHLORIDE	2MG	TABLET;ORAL	Prescription No	AB
HYDROMORPHONE HYDROCHLORIDE	HYDROMORPHONE HYDROCHLORIDE	4MG	TABLET;ORAL	Prescription No	AB
HYDROMORPHONE HYDROCHLORIDE	HYDROMORPHONE HYDROCHLORIDE	8MG	TABLET;ORAL	Prescription No	AB

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

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