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## Drug Details

<b>Drug Name(s)</b>	<b>BUPRENORPHINE HYDROCHLORIDE</b>
<b>FDA Application No.</b>	<b>(ANDA) 090819</b>
<b>Active Ingredient(s)</b>	<b>BUPRENORPHINE HYDROCHLORIDE</b>
<b>Company</b>	<b>ACTAVIS ELIZABETH</b>
<b>Original Approval or Tentative Approval Date</b>	<b>February 19, 2015</b>

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

### Products on Application (ANDA) #090819

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BUPRENORPHINE HYDROCHLORIDE	BUPRENORPHINE HYDROCHLORIDE	EQ 2MG BASE	TABLET;SUBLINGUAL	Prescription No	AB
BUPRENORPHINE HYDROCHLORIDE	BUPRENORPHINE HYDROCHLORIDE	EQ 8MG BASE	TABLET;SUBLINGUAL	Prescription No	AB

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