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## Drug Details

<b>Drug Name(s)</b>	<b>BUSPIRONE HYDROCHLORIDE</b>
<b>FDA Application No.</b>	<b>(ANDA) 204582</b>
<b>Active Ingredient(s)</b>	<b>BUSPIRONE HYDROCHLORIDE</b>
<b>Company</b>	<b>EMCURE PHARMS USA</b>
<b>Original Approval or Tentative Approval Date</b>	<b>September 18, 2015</b>

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

### Products on Application (ANDA) #204582

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

<a href="#">Drug Name</a>	<a href="#">Active Ingredients</a>	<a href="#">Strength</a>	<a href="#">Dosage Form/Route</a>	<a href="#">Marketing Status</a>	<a href="#">RLD</a>	<a href="#">TE Code</a>
BUSPIRONE HYDROCHLORIDE	BUSPIRONE HYDROCHLORIDE	5MG	TABLET;ORAL	Prescription	No	AB
BUSPIRONE HYDROCHLORIDE	BUSPIRONE HYDROCHLORIDE	10MG	TABLET;ORAL	Prescription	No	AB
BUSPIRONE HYDROCHLORIDE	BUSPIRONE HYDROCHLORIDE	15MG	TABLET;ORAL	Prescription	No	AB
BUSPIRONE HYDROCHLORIDE	BUSPIRONE HYDROCHLORIDE	30MG	TABLET;ORAL	Prescription	No	AB

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

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8. <http://www.fda.gov/Drugs/InformationOnDrugs/ucm079436.htm>
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