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## Drug Details

<b>Drug Name(s)</b>	<b>FLUOXETINE HYDROCHLORIDE</b>
<b>FDA Application No.</b>	<b>(ANDA) 206937</b>
<b>Active Ingredient(s)</b>	<b>FLUOXETINE HYDROCHLORIDE</b>
<b>Company</b>	<b>TORRENT PHARMS LTD</b>
<b>Original Approval or Tentative Approval Date</b>	<b>October 21, 2016</b>

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- **Labels are not available**

**Products on Application (ANDA) #206937**  
**Click on a column header to re-sort the table:**

<b>Drug Name</b>	<b>Active Ingredients</b>	<b>Strength</b>	<b>Dosage Form/Route</b>	<b>Marketing Status</b>	<b>RLDTE Code</b>
FLUOXETINE HYDROCHLORIDE	FLUOXETINE HYDROCHLORIDE	EQ 10MG BASE	TABLET;ORAL	Prescription No	AB1
FLUOXETINE HYDROCHLORIDE	FLUOXETINE HYDROCHLORIDE	EQ 20MG BASE	TABLET;ORAL	Prescription No	AB1

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