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## Drug Details

<b>Drug Name(s)</b>	ZOLMITRIPTAN
<b>FDA Application No.</b>	(ANDA) 203772
<b>Active Ingredient(s)</b>	ZOLMITRIPTAN
<b>Company</b>	MACLEODS PHARMS LTD
<b>Original Approval or Tentative Approval Date</b>	September 30, 2015

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

### Products on Application (ANDA) #203772

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>
ZOLMITRIPTAN	ZOLMITRIPTAN	2.5MG	TABLET;ORAL	Prescription	No	AB
ZOLMITRIPTAN	ZOLMITRIPTAN	5MG	TABLET;ORAL	Prescription	No	AB

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