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## Drug Details

<b>Drug Name(s)</b>	<b>RALOXIFENE HYDROCHLORIDE</b>
<b>FDA Application No.</b>	<b>(ANDA) 206384</b>
<b>Active Ingredient(s)</b>	<b>RALOXIFENE HYDROCHLORIDE</b>
<b>Company</b>	<b>SCIEGEN PHARMS INC</b>
<b>Original Approval or Tentative Approval Date</b>	<b>October 12, 2016</b>

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**Products on Application (ANDA) #206384**  
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RALOXIFENE HYDROCHLORIDE	RALOXIFENE HYDROCHLORIDE	60MG	TABLET;ORAL	Prescription	No AB

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