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## Drug Details

<b>Drug Name(s)</b>	<b>ALFUZOSIN HYDROCHLORIDE</b>
<b>FDA Application No.</b>	<b>(ANDA) 203192</b>
<b>Active Ingredient(s)</b>	<b>ALFUZOSIN HYDROCHLORIDE</b>
<b>Company</b>	<b>UNICHEM LABS LTD</b>
<b>Original Approval or Tentative Approval Date</b>	<b>January 28, 2016</b>

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### Products on Application (ANDA) #203192

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ALFUZOSIN HYDROCHLORIDE	ALFUZOSIN HYDROCHLORIDE	10MG	TABLET, EXTENDED RELEASE;ORAL	Prescription No	AB

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