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## Drug Details

<b>Drug Name(s)</b>	<b>MONTELUKAST SODIUM</b>
<b>FDA Application No.</b>	<b>(ANDA) 205695</b>
<b>Active Ingredient(s)</b>	<b>MONTELUKAST SODIUM</b>
<b>Company</b>	<b>ANBISON LAB CO LTD</b>
<b>Original Approval or Tentative Approval Date</b>	<b>November 5, 2015</b>

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

### Products on Application (ANDA) #205695

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<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 TE Code</a>
MONTELUKAST SODIUM	MONTELUKAST SODIUM	EQ 4MG BASE	TABLET, CHEWABLE;ORAL	Prescription	No AB
MONTELUKAST SODIUM	MONTELUKAST SODIUM	EQ 5MG BASE	TABLET, CHEWABLE;ORAL	Prescription	No AB

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