Drug Details

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

- **Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Labels are not available**

Products on Application (ANDA) #205695
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
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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