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Drug Details

Drug Name(s)	MONTELUKAST SODIUM
FDA Application No.	(ANDA) 205683
Active Ingredient(s)	MONTELUKAST SODIUM
Company	ANBISON LAB CO LTD
Original Approval or Tentative Approval Date	January 12, 2016

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

Products on Application (ANDA) #205683

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 10MG BASE	TABLET;ORAL	Prescription	No AB

[Back to Top](#) | [Back to Previous Page](#) | [Back to Drugs@FDA Home](#)

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