Drugs@FDA
FDA Approved Drug Products

Drug Details

Drug Name(s)	**DUTREBIS**
FDA Application No.	**(NDA) 206510**
Active Ingredient(s)	**LAMIVUDINE; RALTEGRAVIR**
Company	**MERCK SHARP DOHME**
Original Approval or Tentative Approval Date	**February 6, 2015**
Chemical Type	**4 New combination**

- **There are no Therapeutic Equivalents**
- **Label Information**
- **Approval History, Letters, Reviews, and Related Documents**

Products on Application (NDA) #206510
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
DUTREBIS	LAMIVUDINE; RALTEGRAVIR	150MG; 300MG	TABLET;ORAL	Prescription	TBD[11]	TBD[12]

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