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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](#)
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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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