Drug Details

Drug Name(s)	**DEXILANT SOLUTAB**
FDA Application No.	**(NDA) 208056**
Active Ingredient(s)	**DEXLANSOPRAZOLE**
Company	**TAKEDA PHARMS USA**
Original Approval or Tentative Approval Date	**January 26, 2016**
Chemical Type	**3 New dosage form**
Review Classification	**S Standard review drug**

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

Products on Application (NDA) #208056
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
DEXILANT SOLUTAB	DEXLANSOPRAZOLE	30MG	TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL	Prescription	Yes	None

Back to Top | Back to Previous Page | Back to Drugs@FDA Home

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