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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](#)
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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	RLDTE Code
DEXILANT SOLUTAB	DEXLANSOPRAZOLE	30MG	TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE; ORAL	Prescription	Yes None

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