Drugs@FDA
FDA Approved Drug Products

Drug Details

Drug Name(s)	**DESVENLAFAXINE SUCCINATE**
FDA Application No.	**(ANDA) 204172**
Active Ingredient(s)	**DESVENLAFAXINE SUCCINATE**
Company	**LUPIN LTD**
Original Approval or Tentative Approval Date	**June 29, 2015**

- **Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Labels are not available**

Products on Application (ANDA) #204172
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
DESVENLAFAXINE SUCCINATE	DESVENLAFAXINE SUCCINATE	EQ 50MG BASE	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB
DESVENLAFAXINE SUCCINATE	DESVENLAFAXINE SUCCINATE	EQ 100MG BASE	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB

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