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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

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- **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	RLDTE 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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