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Drug Details

Drug Name(s)	ESCITALOPRAM OXALATE
FDA Application No.	(ANDA) 202754
Active Ingredient(s)	ESCITALOPRAM OXALATE
Company	MACLEODS PHARMS LTD
Original Approval or Tentative Approval Date	March 31, 2016

- [Therapeutic Equivalents](#)
- [Approval History, Letters, Reviews, and Related Documents](#)
- **Labels are not available**

Products on Application (ANDA) #202754

Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD TE Code
ESCITALOPRAM OXALATE	ESCITALOPRAM OXALATE	EQ 5MG BASE/5ML	SOLUTION;ORAL	Prescription	No AA

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