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Drug Details

Drug Name(s)	QUETIAPINE FUMARATE
FDA Application No.	(ANDA) 203359
Active Ingredient(s)	QUETIAPINE FUMARATE
Company	MACLEODS PHARMS LTD
Original Approval or Tentative Approval Date	May 17, 2016

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

Products on Application (ANDA) #203359

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD TE Code
QUETIAPINE FUMARATE	QUETIAPINE FUMARATE	EQ 25MG BASE	TABLET;ORAL	Prescription	No AB
QUETIAPINE FUMARATE	QUETIAPINE FUMARATE	EQ 50MG BASE	TABLET;ORAL	Prescription	No AB
QUETIAPINE FUMARATE	QUETIAPINE FUMARATE	EQ 100MG BASE	TABLET;ORAL	Prescription	No AB
QUETIAPINE FUMARATE	QUETIAPINE FUMARATE	EQ 200MG BASE	TABLET;ORAL	Prescription	No AB
QUETIAPINE FUMARATE	QUETIAPINE FUMARATE	EQ 300MG BASE	TABLET;ORAL	Prescription	No AB
QUETIAPINE FUMARATE	QUETIAPINE FUMARATE	EQ 400MG BASE	TABLET;ORAL	Prescription	No AB

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