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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