Drug Details

Drug Name(s)	**QUETIAPINE FUMARATE**
FDA Application No.	**(ANDA) 202939**
Active Ingredient(s)	**QUETIAPINE FUMARATE**
Company	**INTELLIPHARMACEUTICS**

- **There are no Therapeutic Equivalents**
- **Labels are not available**
- **Approval History, Letters, Reviews, and Related Documents**

Products on Application (ANDA) #202939
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	50MG	TABLET, EXTENDED RELEASE;ORAL	None (Tentative Approval)	No	None
QUETIAPINE FUMARATE	QUETIAPINE FUMARATE	150MG	TABLET, EXTENDED RELEASE;ORAL	None (Tentative Approval)	No	None
QUETIAPINE FUMARATE	QUETIAPINE FUMARATE	200MG	TABLET, EXTENDED RELEASE;ORAL	None (Tentative Approval)	No	None
QUETIAPINE FUMARATE	QUETIAPINE FUMARATE	300MG	TABLET, EXTENDED RELEASE;ORAL	None (Tentative Approval)	No	None
QUETIAPINE FUMARATE	QUETIAPINE FUMARATE	400MG	TABLET, EXTENDED RELEASE;ORAL	None (Tentative Approval)	No	None

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