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Drug Details

Drug Name(s)	FLUOXETINE HYDROCHLORIDE
FDA Application No.	(ANDA) 204597
Active Ingredient(s)	FLUOXETINE HYDROCHLORIDE
Company	SCIEGEN PHARMS INC
Original Approval or Tentative Approval Date	March 16, 2015

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- **Labels are not available**

Products on Application (ANDA) #204597

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
FLUOXETINE HYDROCHLORIDE	FLUOXETINE HYDROCHLORIDE	EQ 10MG BASE	CAPSULE;ORAL	Prescription No	AB1
FLUOXETINE HYDROCHLORIDE	FLUOXETINE HYDROCHLORIDE	EQ 20MG BASE	CAPSULE;ORAL	Prescription No	AB1
FLUOXETINE HYDROCHLORIDE	FLUOXETINE HYDROCHLORIDE	EQ 40MG BASE	CAPSULE;ORAL	Prescription No	AB

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