Drug Details

Drug Name(s)	**DEXMEDETOMIDINE HYDROCHLORIDE**
FDA Application No.	**(ANDA) 201072**
Active Ingredient(s)	**DEXMEDETOMIDINE HYDROCHLORIDE**
Company	**FRESENIUS KABI USA**
Original Approval or Tentative Approval Date	**September 18, 2015**

- **Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Labels are not available**

Products on Application (ANDA) #201072
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
DEXMEDETOMIDINE HYDROCHLORIDE	DEXMEDETOMIDINE HYDROCHLORIDE	EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)	INJECTABLE;INJECTION	Prescription	No	AP

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