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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	RLDTE Code
DEXMEDETOMIDINE HYDROCHLORIDE	DEXMEDETOMIDINE HYDROCHLORIDE	EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)	INJECTABLE;INJECTION	Prescription No	AP

[Back to Top](#) | [Back to Previous Page](#) | [Back to Drugs@FDA Home](#)

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