Drugs@FDA
FDA Approved Drug Products

Drug Details

Drug Name(s)	**DANTROLENE SODIUM**
FDA Application No.	**(ANDA) 205239**
Active Ingredient(s)	**DANTROLENE SODIUM**
Company	**MYLAN INSTITUTIONAL**
Original Approval or Tentative Approval Date	**February 18, 2016**

- **Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Label Information**

Products on Application (ANDA) #205239
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
DANTROLENE SODIUM	DANTROLENE SODIUM	20MG/VIAL	INJECTABLE;INJECTION	Prescription	No	AP

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