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## Drug Details

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

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- [Approval History, Letters, Reviews, and Related Documents](#)

### Products on Application (ANDA) #205239

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DANTROLENE SODIUM	DANTROLENE SODIUM	20MG/VIAL	INJECTABLE;INJECTION	Prescription	No AP

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