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Drug Details

Drug Name(s)	EVOMELA
FDA Application No.	(NDA) 207155
Active Ingredient(s)	MELPHALAN
Company	SPECTRUM PHARMS
Original Approval or Tentative Approval Date	March 10, 2016
Chemical Type	5 New formulation or new manufacturer

- **There are no Therapeutic Equivalents**
- **Labels are not available**
- [Approval History, Letters, Reviews, and Related Documents](#)

Products on Application (NDA) #207155

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
EVOMELA	MELPHALAN	50MG	INJECTABLE;INJECTION	Prescription	TBD  ¹¹	None

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