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Drug Details

Drug Name(s)	MELPHALAN HYDROCHLORIDE
FDA Application No.	(ANDA) 204773
Active Ingredient(s)	MELPHALAN HYDROCHLORIDE
Company	PAR STERILE PRODUCTS
Original Approval or Tentative Approval Date	August 22, 2016

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- [Approval History, Letters, Reviews, and Related Documents](#)
- **Labels are not available**

Products on Application (ANDA) #204773

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
MELPHALAN HYDROCHLORIDE	MELPHALAN HYDROCHLORIDE	EQ 50MG BASE/VIAL	INJECTABLE;INJECTION	Prescription No	AP

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