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Drug Details

Drug Name(s)	BUSULFAN
FDA Application No.	(ANDA) 202259
Active Ingredient(s)	BUSULFAN
Company	PHARMAFORCE
Original Approval or Tentative Approval Date	December 22, 2015

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

Products on Application (ANDA) #202259

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
BUSULFAN	BUSULFAN	6MG/ML	INJECTABLE;INJECTION	Prescription	No	AP

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