Drug Details

Drug Name(s)	**METHYLPREDNISOLONE SODIUM SUCCINATE**
FDA Application No.	**(ANDA) 202691**
Active Ingredient(s)	**METHYLPREDNISOLONE SODIUM SUCCINATE**
Company	**HIKMA FARMACEUTICA**
Original Approval or Tentative Approval Date	**February 17, 2016**

- **Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Labels are not available**

Products on Application (ANDA) #202691
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
METHYLPREDNISOLONE SODIUM SUCCINATE	METHYLPREDNISOLONE SODIUM SUCCINATE	EQ 500MG BASE/VIAL	INJECTABLE;INJECTION	Prescription	No	AP
METHYLPREDNISOLONE SODIUM SUCCINATE	METHYLPREDNISOLONE SODIUM SUCCINATE	EQ 1GM BASE/VIAL	INJECTABLE;INJECTION	Prescription	No	AP

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