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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

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- [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	RLDTE 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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U.S. Food and Drug Administration
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Silver Spring, MD 20993
Ph. 1-888-INFO-FDA (1-888-463-6332)

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