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Product Details for NDA 011856

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SOLU-MEDROL (METHYLPREDNISOLONE SODIUM SUCCINATE)
EQ 40MG BASE/VIAL Marketing Status: Prescription

SOLU-MEDROL (METHYLPREDNISOLONE SODIUM SUCCINATE)
EQ 125MG BASE/VIAL Marketing Status: Prescription

SOLU-MEDROL (METHYLPREDNISOLONE SODIUM SUCCINATE)
EQ 500MG BASE/VIAL Marketing Status: Prescription

Active Ingredient: METHYLPREDNISOLONE SODIUM SUCCINATE
Proprietary Name: SOLU-MEDROL
Dosage Form; Route of Administration: INJECTABLE; INJECTION
Strength: EQ 500MG BASE/VIAL
Reference Listed Drug: Yes

Reference Standard: Yes
TE Code: AP
Application Number: N011856
Product Number: 005
Approval Date: Approved Prior to Jan 1, 1982
Applicant Holder Full Name: PHARMACIA AND UPJOHN CO
Marketing Status: Prescription
Patent and Exclusivity Information

SOLU-MEDROL (METHYLPREDNISOLONE SODIUM SUCCINATE)
EQ 1GM BASE/VIAL Marketing Status: Prescription

SOLU-MEDROL (METHYLPREDNISOLONE SODIUM SUCCINATE)
EQ 2GM BASE/VIAL Marketing Status: Prescription

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