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Product Details for NDA 011153

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MEDROL (METHYLPREDNISOLONE) 2MG	Marketing Status: Prescription
MEDROL (METHYLPREDNISOLONE) 4MG	Marketing Status: Prescription
MEDROL (METHYLPREDNISOLONE) 8MG	Marketing Status: Prescription
MEDROL (METHYLPREDNISOLONE) 16MG	Marketing Status: Prescription
Active Ingredient: METHYLPREDNISOLONE Proprietary Name: MEDROL Dosage Form; Route of Administration: TABLET; ORAL Strength: 16MG Reference Listed Drug: Yes Reference Standard: No TE Code: AB Application Number: N011153 Product Number: 003 Approval Date: Approved Prior to Jan 1, 1982 Applicant Holder Full Name: PHARMACIA AND UPJOHN CO Marketing Status: Prescription Patent and Exclusivity Information	

MEDROL (METHYLPREDNISOLONE) 32MG	Marketing Status: Prescription
MEDROL (METHYLPREDNISOLONE) 24MG	Marketing Status: Discontinued

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