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Product Details for ANDA 065110

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CYCLOSPORINE (CYCLOSPORINE) 25MG	Marketing Status: Prescription
<div>Active Ingredient: CYCLOSPORINE Proprietary Name: CYCLOSPORINE Dosage Form; Route of Administration: CAPSULE; ORAL Strength: 25MG Reference Listed Drug: No Reference Standard: No TE Code: AB1 Application Number: A065110 Product Number: 003 Approval Date: Mar 29, 2005 Applicant Holder Full Name: IVAX PHARMACEUTICALS INC SUB TEVA PHARMACEUTICALS USA Marketing Status: Prescription Patent and Exclusivity Information</div>	
CYCLOSPORINE (CYCLOSPORINE) 50MG	Marketing Status: Prescription
<div>Active Ingredient: CYCLOSPORINE Proprietary Name: CYCLOSPORINE Dosage Form; Route of Administration: CAPSULE; ORAL Strength: 50MG Reference Listed Drug: No Reference Standard: No TE Code: AB1 Application Number: A065110 Product Number: 001 Approval Date: Mar 29, 2005 Applicant Holder Full Name: IVAX PHARMACEUTICALS INC SUB TEVA PHARMACEUTICALS USA Marketing Status: Prescription Patent and Exclusivity Information</div>	
CYCLOSPORINE (CYCLOSPORINE) 100MG	Marketing Status: Prescription
<div>Active Ingredient: CYCLOSPORINE Proprietary Name: CYCLOSPORINE Dosage Form; Route of Administration: CAPSULE; ORAL Strength: 100MG Reference Listed Drug: No Reference Standard: No TE Code: AB1 Application Number: A065110 Product Number: 002 Approval Date: Mar 29, 2005 Applicant Holder Full Name: IVAX PHARMACEUTICALS INC SUB TEVA PHARMACEUTICALS USA Marketing Status: Prescription Patent and Exclusivity Information</div>	

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