Drug Details

Drug Name(s)	**RESTASIS**
FDA Application No.	**(NDA) 050790**
Active Ingredient(s)	**CYCLOSPORINE**
Company	**ALLERGAN**
Original Approval or Tentative Approval Date	**December 23, 2002**
Chemical Type	**3 New dosage form**
Review Classification	**P Priority review drug**

- **There are no Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Label Information**

Products on Application (NDA) #050790
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
RESTASIS	CYCLOSPORINE	0.05%	EMULSION;OPHTHALMIC	Prescription	Yes	None

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