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FDA Approved Drug Products

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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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