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Drug Details

Drug Name(s)	LACRISERT
FDA Application No.	(NDA) 018771
Active Ingredient(s)	HYDROXYPROPYL CELLULOSE
Company	ATON
Original Approval or Tentative Approval Date	June 1, 1981
Chemical Type	3 New dosage form
Review Classification	S Standard review drug

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

Products on Application (NDA) #018771
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
LACRISERT	HYDROXYPROPYL CELLULOSE	5MG	INSERT;OPHTHALMIC	Prescription	Yes	None

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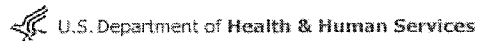
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