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FDA Approved Drug Products

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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](#)
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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

[Back to Top](#) | [Back to Previous Page](#) | [Back to Drugs@FDA Home](#)

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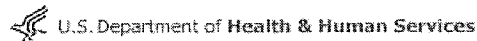
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