TRICARE Prior Authorization Request Form for Ophthalmic Immunomodulatory Agents Subclass: Cyclosporine 0.05% Ophthalmic Emulsion (Restasis)

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JOHNS HOPKINS HEALTHCARE

7231 Parkway Drive, Suite 100, Hanover, MD 21076

FAX Completed Form and Applicable Progress Notes to: (410) 424-4037

USFHP Pharmacy Prior Authorization Form

To be completed by Requesting provider			
Drug Name:	Strength:		
Dosage/Frequency (SIG):	Duration of Therapy:		

Questions? Contact the Pharmacy Dept at: (888) 819-1043, option 4

Clinical Documentation must accompany form in order for a determination to be made.

Step	Please complete patient and physician information (please print):							
1	Patient Name:	Physician Name:						
-	Address:	Address:						
	Sponsor ID #	Phone #:						
O 4	Date of Birth:	Secure Fax #:						
Step	Please complete the clinical assessment:							
2	1. Is this drug being prescribed by an ophthalmologist or optometrist?	☐ Yes	□ No STOP					
		Proceed to question 2	Coverage not approved					
	2. Has the patient received this medication under	□ Yes	□ No					
	the TRICARE benefit in the last 6 months? Please choose "No" if the patient did not previously have a TRICARE approved PA for Restasis	(subject to verification)	Proceed to question 3					
		Proceed to question 12						
	3. Is the patient greater than or equal to 18 years of	□ Yes	□ No					
	age?	Proceed to question 4	STOP					
	4. Will the requested medication be used in	☐ Yes	Coverage not approved No					
	combination with Xiidra or Cequa?	STOP	Proceed to question 5					
		Coverage not approved						
	5. Is the requested medication being prescribed for	□ Yes	□ No					
	LASIK associated dry eyes?	Proceed to question 6	Proceed to question 7					
	6. Did the LASIK surgery occur within the last	□ Yes	□ No					
	THREE Months? Note that therapy is limited to a maximum of THREE months of therapy after the procedure.	Sign and date below	STOP Coverage not approved					
	7. For what indication is the requested medication being prescribed?	□ Moderate to Severe Dry Eye Disease – Proceed to question 8 □ Ocular graft vs. host disease - Sign and date below □ Corneal transplant - Sign and date below □ Atopic keratoconjunctivitis (AKC) - Sign and date below □ Vernal keratoconjunctivitis (VKC) - Sign and date below						
		☐ Other – STOP Coverage not approved						
	8. Has the patient had positive symptomology screening for moderate to severe dry eye disease from an appropriate measure?	☐ Yes Proceed to question 9	☐ No STOP Coverage not approved					

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	 Has the patient had at least one positive diagnostic test (e.g. Tear Film Breakup Time, Osmolarity, Ocular Surface Staining, Schirmer Tear Test)? 	☐ Yes Proceed to question 10	☐ No STOP Coverage not approved
	10. Has the patient tried and failed at least 1 month of one ocular lubricant used at optimal dosing and frequency (e.g. carboxymethylcellulose [Refresh, Celluvisc, Thera Tears, Genteal, etc], polyvinyl alcohol [Liquitears, Refresh Classic, etc], or wetting agents [Systame, Lacrilube)?	☐ Yes Proceed to question 11	□ No STOP Coverage not approved
	11. Has the patient tried and failed at least 1 month of a different ocular lubricant that is non-preserved at optimal dosing and frequency (e.g. carboxymethylcellulose, polyvinyl alcohol, etc.)?	☐ Yes Sign and date below	□ No STOP Coverage not approved
	12. Does the patient have a documented improvement in ocular discomfort?	☐ Yes Proceed to question 13	□ No STOP Coverage not approved
	13. Does the patient have documented improvement in signs of dry eye disease?	☐ Yes Sign and date below	□ No STOP Coverage not approved
	Coverage is not approved for off label uses such as, but not limited to: Pt lens intolerance.	terygia, blepharitis, ocular	
Step 3	I certify the above is true to the best of my knowledge.	Please sign and date	:
	Prescriber Signature	Date	
	<u> </u>		[31 August 20

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For Internal Use Only				
Approved:	Duration of Approval:month(s)			
☐ Denied:	Authorized By:			
☐ Incomplete/Other:	PA#:			