

US Family Health Plan

Prior Authorization Request Form for

Ophthalmic Immunomodulatory Agents Subclass:

Cyclosporine 0.05% Ophthalmic Emulsion (**Restasis**)

To be completed and signed by the prescriber. To be used only for prescriptions which are to be filled through the Department of Defense (DoD) US Family Health Plan Pharmacy Program. US Family Health Plan is a TRICARE contractor for DoD.

The completed form may be **faxed to 855-273-5735**

OR

The patient may attach the completed form to the prescription and mail it to:

Attn: Pharmacy, 77 Warren St, Brighton, MA 02135

QUESTIONS? **Call 1-877-880-7007**

Prior authorization for initial therapy and renewal therapy will approve for 1 time. Consecutive therapy within 120 days will continue without additional Prior authorization required. Therapy requested outside of 120 days will require additional prior authorization. For renewal of therapy an initial Tricare prior authorization approval is required.

usfamilyhealth.org/rx-pa

Step 1 Please complete patient and physician information (please print):

Patient Name: _____ Address: _____ Sponsor ID #: _____ Date of Birth: _____	Physician Name: _____ Address: _____ Phone #: _____ Secure Fax #: _____
--	--

Step 2 Please complete the clinical assessment:

1. Is this drug being prescribed by an ophthalmologist or optometrist?	<input type="checkbox"/> Yes Proceed to question 2	<input type="checkbox"/> No STOP Coverage not approved
2. Has the patient received this medication under the TRICARE benefit in the last 6 months? <i>Please choose "No" if the patient did not previously have a TRICARE approved PA for Restasis</i>	<input type="checkbox"/> Yes (subject to verification) Proceed to question 12	<input type="checkbox"/> No Proceed to question 3
3. Is the patient greater than or equal to 18 years of age?	<input type="checkbox"/> Yes Proceed to question 4	<input type="checkbox"/> No STOP Coverage not approved
4. Will the requested medication be used in combination with Xiidra or Cequa?	<input type="checkbox"/> Yes STOP Coverage not approved	<input type="checkbox"/> No Proceed to question 5
5. Is the requested medication being prescribed for LASIK associated dry eyes?	<input type="checkbox"/> Yes Proceed to question 6	<input type="checkbox"/> No Proceed to question 7
6. Did the LASIK surgery occur within the last THREE Months? <i>Note that therapy is limited to a maximum of THREE months of therapy after the procedure.</i>	<input type="checkbox"/> Yes Sign and date below	<input type="checkbox"/> No STOP Coverage not approved
7. For what indication is the requested medication being prescribed?	<input type="checkbox"/> Moderate to Severe Dry Eye Disease – Proceed to question 8 <input type="checkbox"/> Ocular graft vs. host disease - Sign and date below <input type="checkbox"/> Corneal transplant - Sign and date below <input type="checkbox"/> Atopic keratoconjunctivitis (AKC) - Sign and date below <input type="checkbox"/> Vernal keratoconjunctivitis (VKC) - Sign and date below <input type="checkbox"/> Other – STOP Coverage not approved	
8. Has the patient had positive symptomology screening for moderate to severe dry eye disease from an appropriate measure?	<input type="checkbox"/> Yes Proceed to question 9	<input type="checkbox"/> No STOP Coverage not approved

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

9. Has the patient had at least one positive diagnostic test (e.g. Tear Film Breakup Time, Osmolarity, Ocular Surface Staining, Schirmer Tear Test)?	<input type="checkbox"/> Yes Proceed to question 10	<input type="checkbox"/> 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)?	<input type="checkbox"/> Yes Proceed to question 11	<input type="checkbox"/> 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.)?	<input type="checkbox"/> Yes Sign and date below	<input type="checkbox"/> No STOP Coverage not approved
12. Does the patient have a documented improvement in ocular discomfort?	<input type="checkbox"/> Yes Proceed to question 13	<input type="checkbox"/> No STOP Coverage not approved
13. Does the patient have documented improvement in signs of dry eye disease?	<input type="checkbox"/> Yes Sign and date below	<input type="checkbox"/> No STOP Coverage not approved

Coverage is not approved for off label uses such as, but not limited to: Pterygia, blepharitis, ocular rosacea, and contact lens intolerance.

**Step
3**

I certify the above is true to the best of my knowledge. Please sign and date:

Prescriber Signature

Date

[31 July 2019]