## **HEALTHY U** MEDICAID

## PRIOR AUTHORIZATION REQUEST FORM CABLIVI®

For authorization, please answer each question and fax this form PLUS chart notes back to the Healthy U Prior Authorization Department.

- For **Medical Pharmacy** please fax requests to: 801-213-1547
- For **Retail Pharmacy** please fax requests to: 385-425-4052

Failure to submit clinical documentation to support this request will result in a dismissal of the request.

If you have prior authorization questions, please call for Pharmacy Customer Service for assistance at 385-425-5094

Disclaimer: Prior authorization request forms are subject to change in accordance with Federal and State notice requirements.

| Discialifier. The authorization request for   | a. c sasject to thange in acce | raarice  | Tricir i cuc | state and state fields requirements. |  |
|---|--------------------------------|----------|--------------|--------------------------------------|--|
| Date:   | Member Name:                   |          | ID#:         | ID#:                                 |  |
|   |                                |          |              |                                      |  |
| DOB:  | Gender:                        |          | Phy          | Physician:                           |  |
| Office Phone:   | Office Fax:                    |          | Offi         | Office Contact:                      |  |
|   |                                |          |              | ome contact.                         |  |
| Height/Weight:  |                                |          |              |                                      |  |
| Member must try formulary preferred drugs before a request for a non-preferred drug may be considered. If treatment with preferred products has not been successful, you must submit which preferred products have been tried, dates of treatment, and reason for failure. Reasons for failure must meet the Health Plan medical necessity criteria.  Product being requested:   Cablivi® (caplacizumab-yhdp) |                                |          |              |                                      |  |
| Dosing/Frequency:   |                                |          |              |                                      |  |
| If the request is for reauthorization, proceed to reauthorization section.  |                                |          |              |                                      |  |
| Questions   |                                | Yes      | No           | Comments/Notes                       |  |
| <ol> <li>Does the member have a diagnosis<br/>thrombocytopenia purpura (aTTP) v<br/>&lt;10%?</li> </ol>   | •                              |          |              | Please provide documentation         |  |
| 2. Is the prescriber a hematologist or in consultation with one?  |                                |          |              |                                      |  |
| 3. Is the member 18 years of age or older?  |                                |          |              |                                      |  |
| 4. Will Cablivi® be started in a hospita plasma exchange?   | setting in combination with    |          |              | Please provide documentation         |  |
| 5. Will Cablivi® be used in combination therapy (e.g. corticosteroids, rituxing   | • •                            |          |              | Please provide documentation         |  |
| 6. Have secondary causes of thrombod<br>(e.g. congenital thrombotic thromb<br>hemolytic uremic syndrome, drug-in  | ocytopenia purpura,            |          |              | Please provide documentation         |  |
| REAUTHORIZATION   |                                |          |              |                                      |  |
| 1. Is the request for reauthorization of  | f therapy?                     |          |              |                                      |  |
| 2. Does documentation show persiste an ADAMTS13 activity <20%?  | nt underlying disease with     |          |              | Please provide documentation         |  |
| 3. Has the member experienced >2 reinitial therapy?   | currences of aTTP during       |          |              |                                      |  |
| 4. Has the member demonstrated a possible shown by one of the following:  | ositive response to therapy    |          |              | Please provide documentation         |  |
|   |                                | <u> </u> |              |                                      |  |

| <ul> <li>Clinically significant increase in platelet count (i.e. platelet</li> </ul>                         |  |  |  |  |  |  |  |
|--|--|--|--|--|--|--|--|
| count is within the normal range)  |  |  |  |  |  |  |  |
| <ul> <li>Reduction in neurological symptoms</li> </ul>   |  |  |  |  |  |  |  |
| <ul> <li>Improvement in organ-damage markers (lactate</li> </ul>   |  |  |  |  |  |  |  |
| dehydrogenase, cardiac troponin1 and serum creatinine)   |  |  |  |  |  |  |  |
| What medications and/or treatment modalities have been tried in the past for this condition? Please document |  |  |  |  |  |  |  |
| name of treatment, reason for failure, treatment dates, etc.   |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
| Additional information:  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
| Physician Signature:   |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |

\*\* Failure to submit clinical documentation to support this request will result in a dismissal of the request.\*\*

Policy PHARM-HU-094 Origination Date: 01/01/2022 Reviewed/Revised Date: 05/17/2023 Next Review Date: 05/17/2024 Current Effective Date: 06/01/2023

## **Confidentiality Notice**

This document and any accompanying document contain confidential information and is intended for the use of the individual or entity named on this transmission sheet. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution or the taking of any action in reliance on the contents of this information is strictly prohibited and the document should be returned to this office immediately. If you have received this facsimile in error, please notify us by telephone immediately and destroy document received.