



Cablivi (caplacizumab)

Fax completed form to: (855) 840-1678

If this is an URGENT request, please call (800) 882-4462
(800.88.CIGNA)

PHYSICIAN INFORMATION			PATIENT INFORMATION		
* Physician Name:			*Due to privacy regulations we will not be able to respond via fax with the outcome of our review unless all asterisked (*) items on this form are completed.*		
Specialty:	* DEA, NPI or TIN:				
Office Contact Person:			* Patient Name:		
Office Phone:			* Cigna ID:	* Date of Birth:	
Office Fax:			* Patient Street Address:		
Office Street Address:			City:	State:	Zip:
City:	State:	Zip:	Patient Phone:		
Urgency: <input type="checkbox"/> Standard <input type="checkbox"/> Urgent (In checking this box, I attest to the fact that applying the standard review time frame may seriously jeopardize the customer's life, health, or ability to regain maximum function)					
Medication Requested: <input type="checkbox"/> Cablivi 11mg powder for injection ICD10: Dose & Quantity: Frequency of therapy: Duration of therapy:					
Where will this medication be obtained? <input type="checkbox"/> Biologics <input type="checkbox"/> Hospital Outpatient <input type="checkbox"/> Retail pharmacy <input type="checkbox"/> Other (please specify): <input type="checkbox"/> Home Health / Home Infusion vendor <input type="checkbox"/> Physician's office stock (billing on a medical claim form)					
Facility and/or doctor dispensing and administering medication: Facility Name: State: Tax ID#: Address (City, State, Zip Code):					
Where will this drug be administered? <input type="checkbox"/> Patient's Home <input type="checkbox"/> Hospital Outpatient <input type="checkbox"/> Physician's Office <input type="checkbox"/> Other (please specify): NOTE: Per some Cigna plans, infusion of medication MUST occur in the least intensive, medically appropriate setting.					
Is this patient a candidate for re-direction to an alternate setting (such as alternate infusion site, physician's office, home) with assistance of a Specialty Care Options Case Manager? <input type="checkbox"/> Yes <input type="checkbox"/> No (provide medical necessity rationale):					
Is the requested medication for a chronic or long-term condition for which the prescription medication may be necessary for the life of the patient? <input type="checkbox"/> Yes <input type="checkbox"/> No					
What is your patient's diagnosis? <input type="checkbox"/> acquired (autoimmune) thrombotic thrombocytopenic purpura (aTTP) <input type="checkbox"/> other (please specify):					
Clinical Information Was Cablivi initiated in an inpatient setting to treat acquired (autoimmune) thrombotic thrombocytopenic purpura (aTTP)? <input type="checkbox"/> Yes <input type="checkbox"/> No Was Cablivi initiated in combination with plasma exchange therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No Is the patient currently receiving at least ONE immunosuppressive therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No Is this medication being prescribed by, or in consultation with, a hematologist? <input type="checkbox"/> Yes <input type="checkbox"/> No					

Is this a new start or has the patient previously received Cablivi?

☐ New start

☐ Previously received Cablivi

(if previously received) Has the patient had MORE THAN two (2) recurrences of aTTP while taking Cablivi?

☐ Yes ☐ No

(if new start) When initiated on day 1 of treatment, along with plasma exchange, were two doses of Cablivi given (11 mg intravenous bolus prior to plasma exchange followed by an 11 mg subcutaneous dose after completion of plasma exchange)?

☐ Yes ☐ No

Is the requested dosing 11 mg via subcutaneous injection up to once daily?

☐ Yes ☐ No

Has the patient received over 60 doses of Cablivi following the last plasma exchange session?

☐ Yes ☐ No

Additional pertinent information *(Please provide clinical rationale for the use of this drug for your patient (pertinent patient history, alternatives tried, any inability to use alternatives above or standard therapy, etc). Please include drug name(s), date(s) taken and for how long, and what the documented results were of taking each drug, including any intolerances or adverse reactions your patient experienced.):*

Attestation: I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan or insurer its designees may perform a routine audit and request the medical information necessary to verify the accuracy of the information reported on this form.

Prescriber Signature:_____ **Date:**_____

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