

Tecvayli

Prior Authorization Request

Your patient's benefit plan requires prior authorization for certain medications. In order to make appropriate medical necessity determinations, your patient's diagnosis and other clinical information is required. Please complete the information requested on the form below and fax this form along with supporting clinical documentation to Priority Partners, toll-free at 1-866-212-4756 to initiate the review process. If you have questions regarding the prior authorization please contact Priority Partners at 888-819-1043 Option 4.

Patient's Name:	Date:
Patient's ID:	Patient's Date of Birth:
Physician's Name:	
Specialty:	NPI#:
Physician Office Telephone:	Physician Office Fax:
Referring Provider Info: □ Same as Requesting Pro	ovider
Name:	NPI#:
Fax:	Phone:
Rendering Provider Info: Same as Referring Provider Info:	
Fax:	Phone:
accepted compendia, and/or Required Demographic Information:	r evidence-based practice guidelines.
Patient Weight:kg	
Patient Height:cm	
Please indicate the place of service for the requested dr	ug:
☐ Ambulatory Surgical (POS Code 24)	☐ Home (POS Code 12)
☐ Off Campus Outpatient Hospital (POS Code 19)☐ Office (POS Code 11)	☐ On Campus Outpatient Hospital (POS Code 22)
Drug Information:	
Strength/Measure	Units □ ml □ Gm □ mg □ ea □ Un
	Route of administration
Dosing frequency	
What is the ICD-10 code?	

Note: This fax may contain medical information that is privileged and confidential and is solely for the use of individuals named above. If you are not the intended recipient you hereby are advised that any dissemination, distribution, or copying of this communication is prohibited. If you have received the fax in error, please immediately notify the sender by telephone and destroy the original fax message. JHHC PP Tecvayli SGM 5657-A – 10/2023.

Criteria Questions:	
 1. What is the patient's diagnosis? ☐ Multiple Myeloma, Continue to #2 ☐ Other, Continue to #2 	
<u>Continuation</u>	
 2. Is the patient currently receiving treatment with the requested ☐ Yes, Continue to #3 ☐ No, Continue to #10 	medication?
3. Is there evidence of unacceptable toxicity or disease progression. ☐ Yes, No Further Questions. ☐ No, No Further Questions	on while on the current regimen?
<u>Initial</u>	
10. What is the clinical setting in which the requested medication ☐ Relapsed disease, Continue to #11 ☐ Refractory disease, Continue to #11 ☐ Progressive disease, Continue to #11 ☐ Other, Continue to #11	n will be used?
11. Has the patient received at least four prior therapies for multi- each of the following categories: A) anti-CD38 monoclonal antib- inhibitor (e.g., bortezomib, ixazomib, carfilzomib), and C) immu- pomalidomide)? Yes, No Further Questions No, No Further Questions	oody (e.g., daratumumab), B) proteasome
I attest that this information is accurate and true, and that documents	
information is available for review if requested by Priority Partn	ers.
XPrescriber or Authorized Signature	Date (mm/dd/yy)