Prior Authorization Form



Note: Please provide as much information as possible on this form. Missing data may cause processing delays for requested authorization(s). Attach additional sheets to this form if necessary.

Please fax the completed PA form and any additional informational sheets to Nirvanahealth at the following fax number: +1(866) 871-8565

Patient Information	Prescriber Information	
Patient Name: Health Plan Name: Patient Insurance Id: Patient Date of Birth: Patient Phone:	Prescriber Name: Prescriber Address: Prescriber Phone: () Prescriber Fax: ()	
	Prescriber Specialty: Prescriber DEA: Prescriber NPI:	
Medicat	ation & Medical Information	
Requested Drug(s) & Strength(s):	[] Tarpeyo 4 mg capsule,delayed release	
Requested Daily Quantity Limit – Amount:		
Requested Daily Quantity Limit – Days:		
Expected Length of Therapy:		
Directions:		
Diagnosis and Diagnosis Codes (ICD-10 Standard Codes):		
List drugs used previously to treat the same condition:		
Additional clinical information or history. Please include any relevant test results and/or medical record notes:		
Ougstionnaire		
Questionnaire Q1: I, as the provider or designated representative of the provider, certify and attest that the information provided is complete and accurate and that, upon request, I shall provide any information to RxAdvance that RxAdvance determines is reasonably necessary to verify my responses. (Check only one that apply)		
[] Yes		
[] No		
Q2: What is the member's diagnosis? (Check only one	e that apply)	
[] Primary immunoglobulin A nephropathy (IgAN)		
[] Other (please specify the member's diagnosis and provide clinical rationale for the request)(*Required)		
O2: Is the member at rick of rapid disease progression? (Check only one that apply)		

Q3: Is the member at risk of rapid disease progression? (Check only one that apply)

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[] Yes (please specify)	(*Required)
[] No (please provide clinical rationale for the request)(*Required)	
Q4: Is the requested medication used to reduce proteinuria? (Che	ck only one that apply)
[] Yes	
[] No (please provide clinical rationale for the request)(*Required)	
Q5: Is the estimated glomerular filtration rate (eGFR) greater than	or equal to 35 mL/min/1.73 m2? (Check only one that apply)
[] Yes (please specify)	(*Required)
[] No (please provide clinical rationale for the request)(*Required)	
Q6: Has the member been on a minimum 90-day trial of a maxima of the following? (Check only one that apply)	Illy tolerated dose and will continue to receive therapy with one
[] An angiotensin-converting enzyme (ACE) inhibitor (e.g., bedate of therapy)	nazepril, lisinopril) (please specify drugs name and start and end(*Required)
[] An angiotensin II receptor blocker (ARB) (e.g., losartan, valtherapy)	
[] None of the above	
Q7: Does the member experienced contraindication or intolerance	e to both ACE inhibitors and ARBs? (Check only one that apply)
[] Yes (please specify drug name(s), corresponding contraind (s) of therapy (month/year))	ication(s) or intolerance experienced and the start and end date(*Required)
[] No (please provide clinical rationale for the request) (*Required)	
Q8: Has the member had an inadequate response, intolerance or methylprednisolone, prednisone)? (Check only one that apply)	experienced contraindication(s) to another glucocorticoid (e.g.,
[] Yes (please specify drug name, corresponding contraindicate of therapy (month/year))	ation(s) or intolerance experienced and the start and end date(s)(*Required)
[] No (please provide clinical rationale for the request)(*Required)	
Q9: Is the requested medication prescribed by or in consultation v	vith a nephrologist? (Check only one that apply)
[] Yes	
[] No (please provide clinical rationale for the request)(*Required)	
<u>Attestation:</u> I attest the information provided is true and accurate to the Medical Group or its designated representatives may perform a routine a accuracy of the information reported on this form.	
Signature of Prescriber or Authorized Representative:	Date:
Print Authorized Representative Name:	I