

Kaiser Permanente Health Plan of Mid-Atlantic States, Inc.
Jynarque (tolvaptan) Prior Authorization (PA)
Pharmacy Benefits Prior Authorization Help Desk
Length of Authorizations: Initial- 12 months; Continuation- 12 months

## **Instructions:**

This form is used by Kaiser Permanente and/or participating providers for coverage of **Jynarque (tolvaptan)** for **Commercial, Exchange, FEHB (Federal),** and **MD Medicaid** plans. Please complete and fax this form back to Kaiser Permanente within 24 hours [fax: <u>1-866-331-2104</u>]. If you have any questions or concerns, please call <u>1-866-331-2103</u>. **Requests will not be considered unless this form is complete.** 

KP-MAS Formulary can be found at: <a href="Pharmacy">Pharmacy</a> | Community Provider Portal | Kaiser Permanente</a>

	1 – Patient Information				
Patient Name:	Kaiser Medical ID#:	Date of Birth:			
	2 – Provider Information				
Is the prescriber a nephrologist?   No	□ Yes				
If consulted with a specialist, specialist	name and specialty:				
Provider Name:	Specialty:	NPI:			
Provider Address:					
Provider Phone #:	Provider Fax #:				
Do you have an approved provider refe	erral number from Kaiser Permanente? eferral number here:				
3 — Pharmacy Information					
Pharmacy Name:	Pharmacy NPI:				
Pharmacy Phone #	Pharmacy Fax #:				
	4 – Drug Therapy Requested				
Drug 1: Name/Strength/Formulation: _					
Sig:					
Drug 2: Name/Strength/Formulation: _					
Sig:					

	5— Diagnosis/Clinical Criteria
1.	Is this request for initial or continuing therapy?
	□ Initial therapy □ Continuing therapy, state start date:
2.	Indicate the patient's diagnosis for the requested medication:
Cli	nical Criteria:
1.	Member is ≥18 years and <56 years,  □ No □ Yes
2.	<b>AND</b> eGFR >25 mL/min/1.73 m <sup>2</sup> , $\Box$ No $\Box$ Yes
3.	AND baseline labs completed within 30 days and within normal limits: ALT, AST, bilirubin; and negative pregnancy test (if applicable),  □ No □ Yes
4.	<ul> <li>AND member has a diagnosis of autosomal dominant polycystic kidney disease (ADPKD) confirmed by one of the following:         <ul> <li>Ultrasonography:</li> <li>With family history: ≥3 cysts (unilateral or bilateral) in patients aged 15-39 years OR ≥2 cysts in each kidney in patients aged 40-59 years</li> <li>Without family history: ≥10 cysts per kidney</li> </ul> </li> <li>OR Magnetic resonance imaging (MRI) or computed tomography (CT) scan:         <ul> <li>With family history: ≥5 cysts per kidney</li> <li>Without family history: ≥10 cysts per kidney</li> <li>No □ Yes</li> </ul> </li> </ul>
5.	<ul> <li>AND high risk of disease progression defined by one of the following:         <ul> <li>Mayo ADPKD Classification 1C, 1D, or 1E</li> <li>eGFR decline ≥5 mL/min/1.73 m² in one year OR eGFR decline ≥2.5 mL/min/1.73 m² per year over a period of ≥5 years</li> <li>Truncating PKD1 mutation AND PROPKD score &gt;6</li> <li>No □ Yes</li> </ul> </li> </ul>
	continuation of therapy, please respond to <u>additional questions</u> below. New members who were initiated on therapy side of Kaiser, who have not been reviewed previously, must meet all above Clinical Criteria.
1.	Member has positive clinical response to tolvaptan, No □ Yes
2.	<b>AND</b> member's eGFR >25 mL/min/1.73 m <sup>2</sup> , No $\square$ Yes
3.	AND member has followed up with a nephrologist within the last 12 months No □ Yes

## 7 - Provider Sign-Off

dditional Information –			
. Please submit chart notes/medical records for the	patient that are applicable to this request.		
If member has not tried preferred agent(s) please provide rationale/explanation and any additional supporting information that should be taken into consideration for the requested medication:			
certify that the information provided is accurate. Support	rting documentation is available for State audits		
Provider Signature:	Date:		
Trovider digitature.	Dute.		
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