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This fax machine is located in a secure location as required by HIPAA regulations. Fax complete signed and dated forms to CVS/Caremark at . Please contact CVS/Caremark at 1-888-413-2723 with questions regarding the prior authorization process. When conditions are met, we will authorize the coverage of the medication.

Patient Name: _____ **Date:** 3/16/2026
Patient ID: _____ **Patient Date Of Birth:** _____
Patient Group No: _____ **Patient Phone:** _____ **Physician Name:** _____
 _____ **NPI#:** _____ **Specialty:** _____
 _____ **Physician Office Telephone:** _____

Physician Office Address: _____

Drug Name (specify drug) _____

Quantity: _____ **Frequency:** _____ **Strength:** _____

Route of Administration: _____ **Expected Length of Therapy:** _____

Diagnosis: _____ **ICD Code:** _____

Comments: _____

Please check the appropriate answer for each applicable question.

1. What is the diagnosis?
 - Primary immunoglobulin A nephropathy (IgAN) (If checked, go to 2)
 - Other, please specify (If checked, no further questions)

2. Is the requested drug being prescribed by or in consultation with a nephrologist? Y N

3. Is the patient currently receiving treatment with the requested drug? Y N

4. Has the patient experienced benefit from therapy as evidenced by decreased levels of proteinuria or urine protein-to-creatinine ratio (UPCR) from baseline? ACTION REQUIRED: If Yes, please attach supporting laboratory report or chart note(s). ACTION REQUIRED: Submit supporting documentation Y N

5. Has the diagnosis of primary immunoglobulin A nephropathy (IgAN) been confirmed by a kidney biopsy? ACTION REQUIRED: If Yes, please attach supporting biopsy report confirming diagnosis. ACTION REQUIRED: Submit supporting documentation Y N

6. Has the patient had proteinuria greater than or equal to 0.5 grams per day (g/day) within the last 3 months? ACTION REQUIRED: If Yes, attach supporting laboratory report or chart note(s). ACTION REQUIRED: Submit supporting documentation Y N

7. Has the patient had a urine protein-to-creatinine ratio (UPCR) greater than or equal to 0.8 grams per gram (g/g) within the last 3 months? ACTION REQUIRED: If Yes, please attach supporting laboratory report or chart note(s). ACTION REQUIRED: Submit supporting documentation Y N

8. Has the patient received a stable dose of maximally tolerated renin-angiotensin system (RAS) inhibitor therapy (e.g., angiotensin converting enzyme inhibitor [ACEI] or angiotensin II receptor blocker [ARB]) for at least 3 months prior to initiation of therapy? Y N

9. Does the patient have an intolerance or contraindication to RAS inhibitors? Y N

I attest that the medication requested is medically necessary for this patient. I further attest that the information provided is accurate and true, and that the documentation supporting this information is available for review if requested by the claims processor, the health plan sponsor, or, if applicable a state or federal regulatory agency.

Prescriber (Or Authorized) Signature and Date

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