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**Patient Name:** \_\_\_\_\_ **Date:** 5/13/2025  
**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) ☐

Other, please specify ☐
2. Is the patient currently receiving treatment with the requested medication?
 

Y ☐

N ☐
3. Has the patient experienced benefit from therapy as evidenced by decreased levels of proteinuria or urine protein-to-creatinine ratio (UPCR) from baseline on a 24-hour urine collection? ACTION REQUIRED: If Yes, please attach supporting laboratory report or chart note(s).  
ACTION REQUIRED: Submit supporting documentation
 

Y ☐

N ☐
4. Has the diagnosis of immunoglobulin A nephropathy (IgAN) been confirmed by a kidney biopsy? ACTION REQUIRED: If Yes, please attach supporting chart note(s) or biopsy report supporting diagnosis.  
ACTION REQUIRED: Submit supporting documentation
 

Y ☐

N ☐
5. Does the patient have proteinuria greater than or equal to 1 g/day based on a 24-hour urine collection? ACTION REQUIRED: If Yes, please attach supporting laboratory report or chart note(s).  
ACTION REQUIRED: Submit supporting documentation
 

Y ☐

N ☐
6. Does the patient have a urine protein-to-creatinine ratio (UPCR) greater than or equal to 0.8 g/g based on a 24-hour urine collection? ACTION REQUIRED: If Yes, please attach supporting laboratory report or chart note(s).  
ACTION REQUIRED: Submit supporting documentation
 

Y ☐

N ☐
7. 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 ☐
8. Does the patient have an intolerance or contraindication to RAS inhibitors?
 

Y ☐

N ☐



9. Has the patient experienced an intolerance to oral glucocorticoid (e.g., prednisone)?

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.

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**Prescriber (Or Authorized) Signature and Date**

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