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Patient Name: _____ **Date:** 10/11/2024
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?

Polyneuropathy of hereditary transthyretin-mediated amyloidosis (transthyretin-type familial amyloid polyneuropathy (ATTR-FAP)) (If checked, go to 2)

☐

Other, please specify. (If checked, no further questions)

☐
2. Was the diagnosis confirmed by detection of a mutation in the TTR gene? ACTION REQUIRED: If Yes, attach a copy of testing or analysis confirming a mutation of the TTR gene.

Yes (If checked, go to 3)

☐

No (If checked, no further questions)

☐

Unknown (If checked, no further questions)

☐

ACTION REQUIRED: Submit supporting documentation
3. Does the patient exhibit clinical manifestations of polyneuropathy of hereditary transthyretin-mediated amyloidosis (ATTR-FAP) (e.g., amyloid deposition in biopsy specimens, TTR protein variants in serum, progressive peripheral sensory-motor polyneuropathy)? ACTION REQUIRED: If Yes, attach medical record documentation confirming the patient demonstrates signs and symptoms of polyneuropathy. ACTION REQUIRED: Submit supporting documentation

Y ☐

N ☐
4. Is the patient a liver transplant recipient?

Y ☐

N ☐
5. Will the requested medication be used in combination with patisiran (Onpattro), tafamidis (Vyndaqel, Vyndamax) or vutrisiran (Amvuttra)?

Y ☐

N ☐
6. Is the requested medication prescribed by or in consultation with any of the following: a) neurologist, b) geneticist, or c) physician specializing in the treatment of amyloidosis?

Y ☐

N ☐
7. Is the request for a continuation of therapy with the requested medication?

Y ☐

N ☐
8. Has the patient demonstrated a beneficial response to treatment with the requested drug compared to baseline (e.g., improvement of neuropathy severity and rate of disease progression as demonstrated by the modified Neuropathy Impairment Scale+7 (mNIS+7) composite score, the Norfolk Quality of Life-Diabetic Neuropathy (QoL-DN) total score, polyneuropathy disability (PND) score, FAP disease stage, manual grip strength). ACTION REQUIRED: If Yes, attach chart notes or medical record documentation confirming the clinical benefit of the requested drug. ACTION REQUIRED: Submit supporting documentation

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

Now you can get responses to drug PAs immediately and securely online—without faxes, phone calls, or waiting. How? With electronic prior authorization (ePA)! For more information and to register, go to www.caremark.com/epa.