

**PATIENT INFORMATION**

Patient Name: \_\_\_\_\_ DOB: \_\_\_\_\_ Phone: \_\_\_\_\_ Sex: M F Ht: \_\_\_\_\_ Wt: \_\_\_\_\_ lbs kg  
Primary Language: \_\_\_\_\_ Allergies: \_\_\_\_\_  
Patient Preferred Location: \_\_\_\_\_

**DIAGNOSIS & CLINICAL INFORMATION**

&lt;ICD 10 CODE REQUIRED&gt;

**ICD 10 Code**

- E80.20 Unspecified porphyria
- E80.21 Acute intermittent (hepatic) porphyria
- E80.29 Other porphyria

**REQUIRED:** Demographics & Most Recent: H&P, clinical notes, & medication list. Supporting clinical notes to include any past tried and/or failed therapies, intolerance, outcomes, or contraindications to conventional therapy.

**LAB RESULTS:** Include baseline CMP or AST/ALT and homocysteine level.

**PRESCRIPTION****GIVLAARI (givosiran)****Loading Dose**

- Administer 1.25 mg/kg by subcutaneous injection once for one year
- Administer 2.5mg/kg by subcutaneous injection once for one year

**Referring provider to obtain labs and monitor hepatic function, renal function, and homocysteine as clinically indicated during treatment with Givlaari**

Is the patient on any other disease modifying therapy?    Yes    No  
If yes, please note therapy and last dose: \_\_\_\_\_

**Post Treatment Observations:** The patient is observed for 30 minutes following the first administration.

**Adverse Events:** In the event of an adverse reaction occurring at a Medix Infusion suite, utilize the Medix Infusion adverse reactions protocol.

**Comments:**

\_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

**PRESCRIBER INFORMATION**

Prescriber Name: \_\_\_\_\_ Signature: \_\_\_\_\_  
Date: \_\_\_\_\_ NPI #: \_\_\_\_\_ Specialty: \_\_\_\_\_  
Supervising Physician: \_\_\_\_\_ (If Applicable)  
Address: \_\_\_\_\_ City: \_\_\_\_\_ State: \_\_\_\_\_ Zip: \_\_\_\_\_  
Contact Name: \_\_\_\_\_ Phone: \_\_\_\_\_ Fax: \_\_\_\_\_ Email: \_\_\_\_\_