

PATIENT INFORMATION

Patient Name: _____ DOB: _____ Phone: _____ Sex: M F Ht: _____ Wt: _____ lbs kg

Primary Language: _____ Allergies: _____

Patient Preferred Location: _____

<ICD 10 CODE REQUIRED>

DIAGNOSIS & CLINICAL INFORMATION**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**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: _____