



Updated: 04/2025
DMMA Approved: 04/2025

Request for Prior Authorization for Givlaari (givosiran)

Website Form – www.highmarkhealthoptions.com

Submit request via: Fax - 1-855-476-4158

All requests for Givlaari (givosiran) require a Prior Authorization and will be screened for medical necessity and appropriateness using the criteria listed below.

Givlaari (givosiran) Prior Authorization Criteria:

For all requests for Givlaari (givosiran) all of the following criteria must be met:

Coverage may be provided with a diagnosis of acute hepatic porphyria (AHP) and the following criteria is met:

- Member must be 18 years of age or older
- Must be prescribed by or in consultation with a provider who specializes in porphyria (i.e. hematologist, hepatologist, gastroenterologist)
- Member must have active disease defined as having at least 2 documented porphyria attacks requiring hospitalization, urgent care visits, or IV hemin administration within the last 6 months.
- Documentation the members has had elevated urinary or plasma porphobilinogen (PBG) or aminolevulinic acid (ALA) levels with the past year (reference range must be provided)
- The requested dose and frequency is in accordance with FDA-approved labeling, nationally recognized compendia, and/or evidence-based practice guidelines
- **Initial Duration of Approval:** 6 months
- **Reauthorization criteria**
 - Documentation of a decrease in the number of porphyria attacks that require hospitalization, urgent healthcare visits, or IV hemin administration since starting the medication.
- **Reauthorization Duration of Approval:** 12 months

Coverage may be provided for any non-FDA labeled indication if it is determined that the use is a medically accepted indication supported by nationally recognized pharmacy compendia or peer-reviewed medical literature for treatment of the diagnosis(es) for which it is prescribed. These requests will be reviewed on a case by case basis to determine medical necessity.

Drugs are authorized in generic form unless the branded product is on the preferred drug list or the prescriber has indicated in writing that the branded product is medically necessary. If only the branded product is on the preferred drug list, the generic form will be considered non-preferred and shall not require the prescriber to indicate in writing that the branded product is medically necessary.

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