Geisinger

POLICIES AND PROCEDURE MANUAL

Policy: MBP 277.0

Section: Medical Benefit Pharmaceutical Policy

Subject: Elahere (mirvetuximab soravtansine-gynx)

I. Policy:
Elahere (mirvetuximab soravtansine-gynx)

II. Purpose/Objective:
To provide a policy of coverage regarding Elahere (mirvetuximab soravtansine-gynx)

III. Responsibility:
A. Medical Directors
B. Medical Management
C. Pharmacy Department

IV. Required Definitions
1. Attachment – a supporting document that is developed and maintained by the policy writer or department requiring/authoring the policy.
2. Exhibit – a supporting document developed and maintained in a department other than
3. the department requiring/authoring the policy.
4. Devised – the date the policy was implemented.
5. Revised – the date of every revision to the policy, including typographical and grammatical changes.
6. Reviewed – the date documenting the annual review if the policy has no revisions necessary.

V. Additional Definitions
Medical Necessity or Medically Necessary means Covered Services rendered by a Health Care Provider that the Plan determines are:

a. appropriate for the symptoms and diagnosis or treatment of the Member's condition, illness, disease or injury;
b. provided for the diagnosis and the direct care and treatment of the Member's condition, illness disease or injury;
c. in accordance with current standards good medical treatment practiced by the general medical community;
d. not primarily for the convenience of the Member, or the Member's Health Care Provider; and
e. the most appropriate source or level of service that can safely be provided to the Member. When applied to hospitalization, this further means that the Member requires acute care as an inpatient due to the nature of the services rendered or the Member's condition, and the Member cannot receive safe or adequate care as an outpatient

Medicaid Business Segment
Medically Necessary — A service, item, procedure, or level of care compensable under the Medical Assistance program that is necessary for the proper treatment or management of an illness, injury, or disability is one that:

i. Will, or is reasonably expected to, prevent the onset of an illness, condition, injury or disability.
ii. Will, or is reasonably expected to, reduce or ameliorate the physical, mental or developmental effects of an illness, condition, injury or disability.
iii. Will assist the Member to achieve or maintain maximum functional capacity in performing daily activities, taking into account both the functional capacity of the Member and those functional capacities that are appropriate for Members of the same age.

DESCRIPTION:
Elahere (mirvetuximab soravtansine-gynx) is an antibody-drug conjugate which consists of 3 components, a folate receptor alpha (FRα)-directed monoclonal antibody (IgG1 subtype), a small molecule anti-tubulin agent DM4 (a maytansine derivative), and a linker that covalently attaches DM4 to the mirvetuximab antibody. The antibody portion is a chimeric IgG1 directed against folate receptor alpha (FRα); DM4 is a microtubule inhibitor attached to the antibody via a cleavable linker. Upon binding to FRα, mirvetuximab soravtansine is internalized and then intracellularly releases DM4 via proteolytic cleavage. DM4 disrupts the microtubule network within the cell, resulting in cell cycle arrest and apoptosis.

CRITERIA FOR USE: Requires Prior Authorization by Medical Director or Designee

GRANDFATHER PROVISION – Members already established on therapy are eligible for approval as long as there is medical record documentation that the safety and effectiveness of use for the prescribed indication is supported by Food and Drug Administration (FDA) approval or adequate medical and scientific evidence in the medical literature

Elahere (mirvetuximab soravtansine-gynx) will be considered medically necessary for all lines of business when ALL of the following criteria are met:

- Medical record documentation that Elahere is prescribed by a hematologist or oncologist **AND**
- Medical record documentation of age greater than or equal to 18 years **AND**
- Medical record documentation of a diagnosis of platinum-resistant epithelial ovarian, fallopian tube, or primary peritoneal cancer **AND**
- Medical record documentation of the presence of folate receptor alpha (FRα) tumor expression as determined by an FDA-approved test* **AND**
- Medical record documentation of one to three prior systemic treatment regimens

*The FDA approved test for the measurement of FRα tumor expression is Ventana FOLR1 (FOLR-2.1) RXDX Assay

AUTHORIZATION DURATION: Initial approval will be for **6 months** or less if the reviewing provider feels it is medically appropriate. Subsequent approvals will be for an additional **12 months** or less if the reviewing provider feels it is medically appropriate and will require medical record documentation of continued disease improvement or lack of disease progression. The medication will no longer be covered if patient experiences toxicity or worsening of disease.

Note: For Medicaid (GHP Family), any requests for services that do not meet criteria set in the PARP will be evaluated on a case-by-case basis.

LINE OF BUSINESS:
Eligibility and contract specific benefit limitations and/or exclusions will apply. Coverage statements found in the line of business specific benefit document will supersede this policy.

This policy will be revised as necessary and reviewed no less than annually**.**

Devised: 4/25/23

Revised:

Reviewed: