Drug Policy:

Blenrep™ (belantamab mafodotin-blmf)

I. PURPOSE

To define and describe the accepted indications for Blenrep (belantamab mafodotin-blmf) usage in the treatment of cancer, including FDA approved indications, and off-label indications.

New Century Health (NCH) is responsible for processing all medication requests from network ordering providers. Medications not authorized by NCH may be deemed as not approvable and therefore not reimbursable.

The use of this drug must be supported by one of the following: FDA approved product labeling, CMS-approved compendia, National Comprehensive Cancer Network (NCCN), American Society of Clinical Oncology (ASCO) clinical guidelines, or peer-reviewed literature that meets the requirements of the CMS Medicare Benefit Policy Manual Chapter 15.

II. INDICATIONS FOR USE/INCLUSION CRITERIA

A. PREFERRED MEDICATION GUIDANCE FOR INITIAL REQUEST:

1. When health plan Medicaid coverage provisions—including any applicable PDLs (Preferred Drug Lists)—conflict with the coverage provisions in this drug policy, health plan Medicaid coverage provisions take precedence per the Preferred Drug Guidelines OR

2. When health plan Exchange coverage provisions—including any applicable PDLs (Preferred Drug Lists)—conflict with the coverage provisions in this drug policy, health plan Exchange coverage provisions take precedence per the Preferred Drug Guidelines OR
3. For Health Plans that utilize NCH UM Oncology Clinical Policies as the initial clinical criteria, the Preferred Drug Guidelines shall follow NCH L1 Pathways when applicable, otherwise shall follow NCH drug policies AND

4. Continuation requests of previously approved, non-preferred medication are not subject to this provision AND

5. When available, generic alternatives are preferred over brand-name drugs.

B. Multiple Myeloma
   1. The member has relapsed or refractory multiple myeloma and Blenrep (belantamab mafodotin-blmf) will be used as a single agent AND
   2. The member is refractory to at least 4 prior therapies including an anti-CD38 antibody (e.g. daratumumab), an Immunomodulatory drug (e.g. lenalidomide or pomalidomide), and a proteasome inhibitor (e.g. bortezomib, ixazomib or carfilzomib) AND
   3. Blenrep (belantamab mafodotin-blmf) is supported upon documentation of an ophthalmic exam prior to and following the administrations of Blenrep (belantamab mafodotin-blmf).

III. EXCLUSION CRITERIA
   A. Disease progression on or after treatment with Blenrep (belantamab mafodotin-blmf).
   B. Use of Blenrep (belantamab mafodotin-blmf) in members with ocular toxicities: A baseline (within 3 weeks of initiating therapy) and within 2 weeks prior to each dose, ophthalmic exams are reviewed for ocular adverse reactions.
   C. Dosing exceeds single dose limit of Blenrep (belantamab mafodotin-blmf) 2.5 mg/kg.
   D. Indications not supported by CMS recognized compendia or acceptable peer reviewed literature.

IV. MEDICATION MANAGEMENT
   A. Grade 3 or 4 febrile neutropenia is 0% (low risk level).^A
   B. The frequency of emesis is 15.8% (low risk level).^A
   C. Please refer to the FDA label/package insert for details regarding these topics.

V. APPROVAL AUTHORITY
   A. Review – Utilization Management Department
   B. Final Approval – Utilization Management Committee

VI. ATTACHMENTS
   A. None

VII. REFERENCES
   B. Blenrep PI prescribing information. GlaxoSmithKline LLC, Wilmington, Delaware 2020