

| Field Name | Field Description |
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| Prior Authorization Group Description | Anti-CD19 CAR-T Immunotherapies |
| Drugs | Kymriah (tisagenlecleucel), Yescarta (axicabtagene ciloleucel), Tecartus (brexucabtagene autoleucel), Brevanzi (lisocabtagene maraleucel) |
| Covered Uses | Medically accepted indications are defined using the following sources: the Food and Drug Administration (FDA), Micromedex, American Hospital Formulary Service (AHFS), United States Pharmacopeia Drug Information for the Healthcare Professional (USP DI), the Drug Package Insert (PPI), or disease state specific standard of care guidelines. |
| Exclusion Criteria | Patients with primary central nervous system lymphoma |
| Required Medical Information | See “Other Criteria” |
| Age Restrictions | See “Other Criteria” |
| Prescriber Restrictions | Prescriber must be an oncologist, hematologist or other prescribers who specialize in the treatment of lymphoma. |
| Coverage Duration | If all the criteria are met, the initial request will be approved for a one – time infusion per lifetime. |
| Other Criteria | <p>**Drug is being requested through the member’s medical benefit**</p> <p>Initial authorization:</p> <ul style="list-style-type: none"> • Patient must not have received prior anti-CD19 CAR-T therapy. • Patient will be screened for HBV, HCV, and HIV in accordance with clinical guidelines. • Patient does not have an active infection or inflammatory disorder. • Patient has a life expectancy >12 weeks. • Patient will not receive live virus vaccines for at least 6 weeks prior to the start of lymphodepleting chemotherapy and until immune recovery following treatment. <p>Leukemia</p> <p>B-cell precursor Acute Lymphoblastic Leukemia (ALL):</p> <ul style="list-style-type: none"> • If the request is for Kymriah <ul style="list-style-type: none"> ○ Patient is 25 years of age or younger ○ ALL that is refractory or in second or later relapse • <u>If the request is for Tecartus</u> <ul style="list-style-type: none"> ○ <u>Patient is 18 years of age or older</u> ○ <u>ALL that is relapsed or refractory</u> <p>Non-Hodgkin’s Lymphoma (NHL)</p> <p>Mantle Cell Lymphoma (MCL):</p> |

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| <p>Revision/Review Date: 5/2021 <u>1/2022</u></p> | <ul style="list-style-type: none"> • If the request is for Tecartus: <ul style="list-style-type: none"> ○ Patient is 18 years of age or older ○ Patient has relapsed/refractory disease defined as failure of BOTH the following lines of therapy: <ul style="list-style-type: none"> ▪ Chemoimmunotherapy such as an anti-CD20 monoclonal antibody (e.g. Rituxan) + any chemotherapeutic agent ▪ Bruton Tyrosine Kinase (BTK) Inhibitor (e.g. Calquence, Imbruvica, Brukinsa) <p>Other forms of NHL:</p> <ul style="list-style-type: none"> • If the request is for Breyanzi (lisocabtagene maraleucel), Kymriah (tisagenlecleucel), Yescarta (axicabtagene ciloleucel) <ul style="list-style-type: none"> ○ Use is supported by a labeled indication or NCCN guidelines ○ Patient is 18 years of age or older ○ Patient has relapsed/refractory disease defined as failure of two or more lines of systemic therapy <p>Re-authorization:</p> <ul style="list-style-type: none"> • Treatment exceeding 1 dose per lifetime will not be authorized. <p>Medical Director/clinical reviewer must override criteria when, in his/her professional judgement, the requested item is medically necessary.</p> |
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