Division: Pharmacy Policy	Subject: State of Florida's Agency for Health Care Administration's Prior Authorization Criteria
Original Development Date: Original Effective Date: Revision Date:	December 22, 2017 July 16, 2020, March 17, 2021; June 16, 2022

YESCARTA® (axicabtagene ciloleucel suspension)

LENGTH OF AUTHORIZATION: Date of service

REVIEW CRITERIA:

- Patient must be 18 years of age or older.
- Must have large B-cell lymphoma that is refractory to 1st line chemoimmunotherapy or that relapses within 12 months of 1st line chemoimmunotherapy; **OR**
- Must have relapsed or refractory large B-cell lymphoma after two or more lines of treatment with systemic therapy for diffuse large B-cell lymphoma (DLBCL) not otherwise specified, primary mediastinal large B-cell lymphoma, and DLBCL arising from follicular lymphoma.

DOSING AND ADMINISTRATION:

- Refer to product labeling at https://www.accessdata.fda.gov/scripts/cder/daf/
- Because of the risk of Cytokine Release Syndrome (CRS) and neurological toxicities, YESCARTA ® is available only through a restricted program under a Risk Evaluation and Mitigation Strategy (REMS) called the YESCARTA and TECARTUS REMS program. Further information is available at www.YescartaTecartusREMS.com or 1-844-454-KITE (5483).