



MT2021-11: An Open-label, Single-arm, Multicohort, Phase 2 Study to Assess

the Efficacy and Safety of Tabelecleucel in Subjects with Epstein-Barr Virus-associated Diseases Status: Recruiting

Eligibility Criteria

Sex: Male or Female Age Group: Not specified This study is NOT accepting healthy volunteers

Inclusion Criteria:

- diagnosis of Epstein-Barr Virus (EBV) disorder - able to walk and do all selfcare but unable to carry out any work activities; up and about more than 50% of waking hours - see link to clinicaltrials.com for additional inclusion criteria

Exclusion Criteria:

- women who are breastfeeding or pregnant - currently active Burkitt, T-cell, natural killer/T-cell lymphoma/LPD, Hodgkin, plasmablastic, transformed lymphoma, active hemophagocytic lymphohistiocytosis, or other malignancies requiring systemic therapy - serious known active infections - additional exclusion criteria apply (study staff will review)

Conditions & Interventions

Conditions: Cancer, Infectious Diseases Keywords: Clinics and Surgery Center (CSC), Epstein-Barr Virus (EBV)

More Information

Description: This research is being done to determine whether the investigational drug tabelecleucel (allogeneic Epstein-Barr virus-specific cytotoxic T lymphocytes [EBV-CTLs]) can help people with EBV-associated diseases. Study Contact: Eric Homan - homa0030@umn.edu Principal Investigator: Supriya Gupta Phase: PHASE2 IRB Number: STUDY00013494

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