Molecular profiling of tissue samples from patients who received a Kite-manufactured gene-modified cell therapy and have developed a secondary malignancy of T-cell origin

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Last updated: 05/02/2026

Study Planned

Administrative details

EU PAS number

EUPAS1000000769

Study ID

1000000769

DARWIN EU® study

No

Study countries

- ☐ European Union
- ☐ United Kingdom

Study description

KT-EU-982-0910: This is an observational study of patients who received a Kite-manufactured chimeric antigen receptor (CAR) T-cell therapy (axicabtagene ciloleucel or brexucabtagene) and have reported a secondary malignancy of T-cell origin.

The primary objective of this study is to assess potential chimeric antigen receptor (CAR) transgene involvement by performing molecular profiling of tissue and blood samples obtained or are about to be obtained during clinical evaluation, from patients who were treated with axicabtagene ciloleucel or brexucabtagene autoleucel and are suspected to have developed a secondary T-cell malignancy.

Study status

Planned

Research institutions and networks

Institutions

Gilead Sciences

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Institution | Pharmaceutical company

Kite, a Gilead Company

Contact details

Study institution contact

Kite Study Director ClinicalTrialDisclosure@gilead.com

Study contact

ClinicalTrialDisclosure@gilead.com

Primary lead investigator

Kite Study Director

Primary lead investigator

Study timelines

Date when funding contract was signed

Planned: 08/11/2025

Study start date

Planned: 30/06/2026

Date of final study report

Planned: 10/01/2040

Study protocol

982-0910-appendix-16.1.9-protocol_f-redact-FCLTCL0YM74.pdf (2.11 MB)

Regulatory

Was the study required by a regulatory body?

Yes

Is the study required by a Risk Management Plan (RMP)?

EU RMP category 3 (required)

Methodological aspects

Study type

Study type list

Study topic:

Disease /health condition

Human medicinal product

Study type:

Non-interventional study

Scope of the study:

Feasibility analysis

Study Design

Non-interventional study design

Other

Non-interventional study design, other

Observational study

Study drug and medical condition

Medicinal product name

YESCARTA

TECARTUS

Study drug International non-proprietary name (INN) or common name

AXICABTAGENE CILOLEUCEL

BREXUCABTAGENE AUTOLEUCEL

Medical condition to be studied

Diffuse large B-cell lymphoma

Mantle cell lymphoma

Follicular lymphoma

Acute lymphocytic leukaemia

Primary mediastinal large B-cell lymphoma

Population studied

Age groups

- **Adult and elderly population (≥18 years)**
  - Adults (18 to < 65 years)
    - Adults (18 to < 46 years)
    - Adults (46 to < 65 years)
  - Elderly (≥ 65 years)
    - Adults (65 to < 75 years)

- Adults (75 to < 85 years)
- Adults (85 years and over)

Data management

ENCePP Seal

The use of the ENCePP Seal has been discontinued since February 2025. The ENCePP Seal fields are retained in the display mode for transparency but are no longer maintained.

Use of a Common Data Model (CDM)

CDM mapping

No

Data quality specifications

Check conformance

Yes

Check completeness

Yes

Check stability

Yes

Check logical consistency

Yes

Data characterisation

Data characterisation conducted

Unknown