

A Clinical Study Evaluating the Safety and Efficacy of GT729 Universal Cell Injection in the Treatment of Refractory or Relapsed Chronic Graft-versus-host Disease (cGVHD)

NCT07253259

Status	RECRUITING
Phase	Early Phase 1
Sponsor	Grit Biotechnology
Enrollment	36 participants

Key Eligibility Criteria

Inclusion (12)

- Participants or their legal representatives voluntarily sign a written informed consent form, are willing and able to comply with the procedures of this study.
 - Aged 18 to 65 years old (inclusive) when signing the informed consent, regardless of gender.
 - Participants must meet the following criteria:
 - The subjects are allogeneic hematopoietic stem cell transplantation (alloHSCT) recipients with active chronic graft-versus-host disease (active cGVHD) requiring systemic immunosuppressive therapy.
 - The subjects are patients with refractory or relapsed active cGVHD after receiving at least two lines of systemic treatment.
- ... and 7 more (see full listing online)

Exclusion (10)

- Evidence of recurrence of underlying malignant tumors or post-transplant lymphoproliferative disorder (PTLD) at the time of screening
 - Having a history of severe hypersensitivity or allergies
 - Suffering from the following heart diseases:
 - New York Heart Association (NYHA) Class III or IV congestive heart failure;
 - A myocardial infarction occurred or coronary artery bypass surgery was performed within 6 months before the screening period.
- ... and 5 more (see full listing online)

Locations (1 total)

Institute of Hematology & Blood Diseases Hospital, Chinese Academy of Medical Sciences, Tianjin, Tianjin Municipality, China

<https://clinicaltrials.gov/study/NCT07253259>

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