

# T Cell Therapy of Opportunistic Cytomegalovirus Infection

NCT02982902

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<b>Status</b>	RECRUITING
<b>Phase</b>	Early Phase 1
<b>Sponsor</b>	Mari Dallas
<b>Enrollment</b>	20 participants

## Key Eligibility Criteria

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### Inclusion (11)

- Patients must have received allogeneic hematopoietic stem cell transplant and be greater than 30 days post-transplant at the time of registration
- Patients must have documented opportunistic CMV infection, or reactivation; the criteria include (both of the following criteria must be met)
- Patients may have asymptomatic viremia ( $\geq 1000$  copies/ml) OR presence of symptoms secondary to CMV infection, AND
- Patients must have ONE OF THE NEXT FOUR CRITERIA:
- Absence of an improvement of viral load after 14 days of antiviral therapy with ganciclovir, valganciclovir or foscarnet (decrease by at least 1 log, i.e. 10-fold) or
- ... and 6 more (see full listing online)

### Exclusion (11)

- Pregnant or breastfeeding women are excluded from this study.
- Patients with opportunistic viral infections other than CMV.
- Patients with active, grade 2-4, acute graft vs. host disease (GVHD), chronic GVHD or any condition requiring high doses of glucocorticosteroid ( $>0.5$  mg/kg/day prednisone or its equivalent) as treatment
- Treatment with antithymocyte globulin within 28 days of planned infusion of virus - specific, antigen selected T cells.
- Treatment with virus - specific T cells within 6 weeks (42 days) of planned infusion.
- ... and 6 more (see full listing online)

## Locations (1 total)

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University Hospitals Cleveland Medical Center, Case Comprehensive Cancer Center, Cleveland, Ohio, United States

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<https://clinicaltrials.gov/study/NCT02982902>

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