

Human-induced pluripotent stem cell-derived glial enriched progenitors to treat white matter stroke and vascular dementia.

### Grant Award Details

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Human-induced pluripotent stem cell-derived glial enriched progenitors to treat white matter stroke and vascular dementia.

**Grant Type:** Quest - Discovery Stage Research Projects

**Grant Number:** DISC2-12169

**Project Objective:** To complete tumorigenicity, stability and assay development studies, to declare an hiPSC candidate ready for translational stage activities.

**Investigator:**

<b>Name:</b>	Stanley Carmichael
<b>Institution:</b>	University of California, Los Angeles
<b>Type:</b>	PI

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**Disease Focus:** Neurological Disorders, Stroke

**Human Stem Cell Use:** iPS Cell

**Award Value:** \$250,000

**Status:** Pre-Active

### Grant Application Details

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**Application Title:** Human-induced pluripotent stem cell-derived glial enriched progenitors to treat white matter stroke and vascular dementia.

**Public Abstract:** **Research Objective**

This grant proposes development of a stem cell based therapy that is derived from human induced pluripotent stem cells. These cells are in the form of a brain support cell, an astrocyte.

**Impact**

The cell candidate will treat vascular dementia, the second leading cause of dementia, and stroke by overcoming a bottleneck in the ability to make large quantities of the cells for clinical use.

**Major Proposed Activities**

- In vivo tumorigenic studies.
- Development and optimization of potency assays
- Qualification and stability of cell delivery system.

**Statement of Benefit to California:** This research will develop a therapy for a disease with no treatment, vascular dementia, that is common and devastating in its consequences. The intellectual property for this therapy is held by a State of California public university (UCLA) and commercialization will directly benefit the State of California.

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