Microglia based Alzheimer therapy

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Alzheimer’s disease (AD) is an age-related neurodegenerative disease associated with the formation of amyloid plaques, tau aggregation and oxidative/inflammatory damage. Microglia play an important role in the early phase of the disease and are known to be involved in AD progression. Senescent microglia accumulate in AD causing inflammation, neuronal damage and increasing the Aβ load due to a failing protein degradation system.

This study examines the transplantation of in vitro derived young microglia from wild type mice (aged ~3 months) to the brain of AD mice (APP/PS1 transgenic model: >12 months).

Microglia could be detected in the brain for four weeks after transplantation. Analysis of amyloid content shows a significant reduction as well as improved inflammation markers. Microglia and astrocyte morphology and activation status was analysed using histology. Changes in the short term memory of the recipient animals was analysed using a maze and found to be improved. Current analysis focuses on the quality and M1/M2 status of the microglia produced for transplantation to better understand if the therapeutic effect can be improved.

Keywords: Alzheimer, microglia, aging, inflammation