

# Developing cognitive and neurocognitive biomarkers for recovery from alcohol dependence (AD)

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Alcohol is the 7th leading risk for death and disability [1], and there are an estimated 586,780 dependent drinkers in UK [2].



Treatment should be tailored and consider individual-level and structural determinants [3].



Is Brain Gauge a device that could accomplish this, identify patients at risk of relapse, and complement treatment plans?



Currently, formal comprehensive assessment or tracking of this is not used widely in AD treatment [12], though it could be useful when creating treatment plans.



AD linked to impaired prefrontal cortex [4, 5, 6] and therefore impaired executive function [7, 8]. These impairments predict treatment outcomes [9, 10, 11].



Brain Gauge stimulates two adjacent fingertips [13]...



...which creates predictable reactions in adjacent areas of the cortex, affecting perception. Responses give insight into the state of the relevant mechanism.



Brain Gauge has successfully tracked cognitive recovery after mild brain injury [13].



We are now undertaking a longer study with a larger sample to find out more, and to learn whether Brain Gauge has predictive utility.



Our pilot study found that Brain Gauge was able to highlight certain changes across early AD recovery [15].



And has been able to highlight differences in chronic alcohol use against normative data [14].

## Planned Methods

N = 90 AD, 90 one-time controls  
 5 time-points (up to 5-7 months)

### Measures:

- Brain Gauge
- EFI
- HADS
- AUDIT, SADQ, TLFB-A
- SURE

## Planned Analyses

### Within-groups:

RM analyses assessing EFI and Brain Gauge. Regression assessing predictability on recovery indicators

### Between-groups:

Independent analyses comparing scores between AD and controls

## References

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