



Using wearable mHealth devices to maximise outcomes in young people with mental illness: The unWIRED project Anthony Harris^{1,2,3}, Rachael Foord¹, David Johnston¹, Achim Casties⁴, Gordon McDonald⁴, & Kristian Maras⁴



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Background

Community mental health services with an orientation to early intervention strive to intervene at the earliest opportunity in the onset of mental illness or relapse. However, community-based clinicians remain at a disadvantage as they 'catch up' with the mental state of

Results

- 11, male = 8). Average age = 21 years (SD = 1.86 years), all recruited from a young adults' early intervention service, within community mental health WSLHD
- Participants were enrolled in the study for an average of 92.8 days (SD = 44.30 days).
- their clients often *after* a deterioration in a person's mental state, leading to periods of untreated illness [1,2] mHealth (e.g. digital apps and wearables) provide the possibility of real time tracking of key psychophysiological indices of good mental health such as sleep, arousal and activity [3].
- The unWIRED Project integrates a mHealth device into clinical care to track and record these indices with the aim of recognising signs of early deterioration in young adults' mental health.

Aims

- To assess the effectiveness of integrating a mHealth device in the clinical care of young people with a mental illness
- Examine the use of a machine learning approach in predicting deterioration in participants using

- A Random Forest Classifier was superior to a Logistic Classifier; we were able to predict clinical deterioration with a high degree of specificity (0.96) but a lower degree of sensitivity (0.50). Accuracy rate = 0.86 (95% CI 0.75 -0.94).
- SMS self-reported stress levels (via EMA) did not improve the accuracy of the model





Qualitative data suggests that the clients and clinicians identify the benefits of the mHealth device in aiding both self-monitoring and clinician-monitoring [4]

psychophysiological data collected by the mHealth device.



Method

An initial pilot study of the use of the E2 (Empatica) device measuring sleep, arousal (electrodermal activity; EDA), and level of activity was conducted in young adults referred to a community mental health service.

The barrier to utilisation, understandably, is concerns with privacy and data misuse [4]

Conclusion

- We have been able to establish the acceptability of wearing a mHealth device over an extended period
- While reasonable accuracy was achieved it did not reach an acceptable level for clinical use.
- EMA vis SMS self-reporting of stress does appear necessary in monitoring patient deterioration when other physophysiological and clinical data is available
- A subsequent Randomised Controlled Trial (N = 80) is ongoing in 2022 with the aim to further strengthen these findings.

- Labels for participants' functioning ('stable', 'ambivalent' or 'distress') were extracted from medical records and matched with electrophysiological data.
- Ecological Momentary Assessment (EMA) in the form of replies to randomly sent text messages to participants asking them to rate their stress levels on a 1-10 scale were also included.
- Data was examined using machine learning to predict significant changes in mental state.
- Qualitative data from participants and clinician interviews assessed acceptability of the device.

References

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This study has been made possible through the generosity of The Balnaves Foundation

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