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contact with active researchers in the field. The selected publications included both single and multi-centre studies. The Hyperfine Swoop system was used in almost all studies. Mean participant age range was 31 to 63. Qualitative and quantitative comparisons demonstrated good correspondence between high field and ULF-MRI across a range of measures studied, including volumetric measures and moderate to severe white matter hyperintensities.

Conclusion. The limited available evidence suggests that there is potential for ULF-MRI to transform the approach to neuroimaging in the assessment of dementia. Dedicated research into the use of ULF-MRI in this specific application will determine if it will be one of the much-needed disruptors to our current processes of dementia assessment.

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Youth Group Wellbeing Project for Adolescents Impacted by the March 15 Attacks – Protocol for a Pilot Randomised Waitlist-Controlled Trial

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Aims. Psychological distress is common in adolescence; even more so following traumatic events. On 15 March 2019, two mosques in Christchurch were targeted in an act of terrorism. This has had widespread repercussions in the Muslim and wider community in Christchurch and New Zealand. This protocol offers an integrated group treatment based on an indigenous Islamic Psychology framework incorporating components of established transdiagnostic interventions for increasing wellbeing and reducing psychological distress in teenagers. We aim to measure the effect size of the treatment effect on total difficulties, emotional difficulties, trauma symptoms, somatic symptoms and functional impairment in adolescents self-identifying as impacted by the terror attack. We will measure the degree of parental distress and somatic symptoms to explore whether an intervention for adolescents has an impact on parental wellbeing. We will determine the feasibility and acceptability of this approach to inform supports for similar populations and as an example of cultural adaptation of mental health services.

Methods. This is a randomised controlled trial with a waitlistcontrolled design to measure the size of treatment effects on clinical outcomes, and the feasibility of this protocol. We aim to recruit 64 participant families. A 6-week group programme will be offered to teenage participants randomised to the study group and offered to the waitlist group following the study. The study will be community-based in one site. We will assess clinical outcomes including emotional difficulties and somatic symptoms in teenagers (aged 12-19) and parents at baseline, end of treatment and at 3-month follow-up, and measure the project's acceptability with participants and parents. Individuals' experiences of the programme will be examined using qualitative analysis of participant interviews at the end of the programme. Statistical analysis will be a mixed method design including effect size difference calculations, quantitative measures of acceptability and qualitative analysis. Treatment data from participants randomised to the waitlist first will not be included in statistical

comparison of treatment effects but will be used for the assessment of feasibility.

Results. This study will inform whether this unique approach is feasible and easily accessible for adolescents impacted by traumatic events. Its design has been driven by community engagement and stakeholder consultation to consider recruitment, relational safety, screening, and risk management. The project has an emphasis on widening access to mental health supports in a minority faith community by maintaining cultural sensitivity and reducing stigma associated with mental illness.

Conclusion.

Trial registration: ClinicalTrials.gov, NCT05030909. Registered on 8 September 2021.

Abstracts were reviewed by the RCPsych Academic Faculty rather than by the standard *BJPsych Open* peer review process and should not be quoted as peer-reviewed by *BJPsych Open* in any subsequent publication.

Exploring Pathways to Autism Spectrum Diagnosis in Adulthood

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Aims. Autism spectrum is a neurodevelopmental condition usually diagnosed in early childhood. The broadened diagnostic criteria of the DSM-5 (2013) have led to an increasing number of autism spectrum diagnoses of individuals requiring lower levels of support. Barriers to diagnosis, especially in adults, include the complexity of differential diagnosis with co-occurring psychiatric disorders. This study explored the various pathways of psychiatric diagnosis preceding an autism spectrum diagnosis in adulthood.

Methods. This retrospective cohort study was extracted from health-administrative data from Quebec (Canada) and included all adults with a first recorded autism spectrum diagnosis between 2010 and 2017 (index date). A Trajectory of psychiatric Diagnoses (TDx) was defined as a succession of categorical states, each corresponding to a medical record of a psychiatric diagnosis. These TDx were analysed from 2002 to 2017, using a state sequence analysis with trimester as time units. For each trimester, we defined the following diagnoses in order of priority: 1) autism spectrum, 2) intellectual disability (ID), 3) schizophrenia, 4) bipolar disorder (BP), 5) depressive disorder (DD), 6) anxiety disorder (AD), 7) attention-deficit/hyperactivity disorder (ADHD), and 8) other psychiatric disorders. The simple Hamming metric was used to measure the dissimilarity between TDx, followed by a hierarchical cluster analysis to categorise similar trajectories.

Results. The study cohort included 2799 adults diagnosed with autism spectrum between 2010 and 2017. Several psychiatric disorders were recorded during the study period, including AD (77.5%), DD (58.0%), schizophrenia (49.4%), BP (48.3%) and ID (33.2%). Results revealed 5 distinct types of TDx. Types 1 and 2, shared by 63.8% and 17.6% of the cohort respectively, represented individuals in younger age groups with similar characteristics, but with very different sequences of psychiatric diagnoses. Slight or sharp increases in diagnoses were observed around 2010, predominantly associated to autism spectrum in Type 1, and to schizophrenia and AD in Type 2. Individuals