Acceptance and commitment therapy for young brain tumour survivors: study protocol for an acceptability and feasibility trial

Sam Malins, Ray Owen, Ingram Wright, Heather Borrill, Jenny Limond, Faith Gibson, Richard G Grundy, Simon Bailey, Steven C Clifford, Stephen Lowis, James Lemon, Louise Hayes, Sophie Thomas

ABSTRACT

Introduction Survivors of childhood brain tumours have the poorest health-related quality of life of all cancer survivors due to the multiple physical and psychological sequelae of brain tumours and their treatment. Remotely delivered acceptance and commitment therapy (ACT) may be a suitable and accessible psychological intervention to support young people who have survived brain tumours. This study aims to assess the feasibility and acceptability of remotely delivered ACT to improve quality of life among these young survivors.

Methods and analysis This study is a two-arm, parallel group, randomised controlled trial comparing ACT with waitlist control at 12-week follow-up as the primary endpoint. Seventy-two participants will be recruited, who are aged 11–24 and have completed brain tumour treatment. Participants will be randomised to receive 12 weeks of ACT either immediately or after a 12-week wait. The DNA-v model of ACT will be employed, which is a developmentally appropriate model for young people. Feasibility will be assessed using the proportion of those showing interest who consent to the trial and complete the intervention. Acceptability will be assessed using participant evaluations of the intervention, alongside qualitative interviews and treatment diaries analysed thematically. A range of clinical outcome measures will also assess physical and mental health, everyday functioning, quality of life and service usage at 12-week follow-up. The duration of treatment effects will be assessed by further follow-up assessments at 24 weeks, 36 weeks and 48 weeks.

Ethics and dissemination Ethical approval was given by East Midlands, Nottingham 1 Research Ethics Committee (Reference: 20/EM/0237). Study results will be disseminated in peer-reviewed journals, through public events and relevant third sector organisations.

INTRODUCTION

Due to treatment advances, childhood brain tumour survival rates continue to improve, but survival is often associated with significant morbidity. Physical symptoms such as fatigue, and cognitive deficits such as memory difficulties are commonly coupled with psychological problems such as anxiety and post-traumatic stress. Consequently, people who survive childhood brain tumours are identified as having the poorest quality of life of all cancer survivors. In the context of the COVID-19 pandemic, disrupted education, social isolation, financial loss to parents/carers and direct COVID-19-related ill-health also has an especially severe impact on young brain tumour survivors, due to pre-existing vulnerability.

The identification of a psychological care package to improve well-being, social functioning and mental health is regarded as the top research priority by young people who have experienced cancer and their carers. Despite this, there are few studies of psychological therapies for brain tumour patients and none spanning the developmentally important transition from childhood to adulthood.

Acceptance and commitment therapy (ACT) is an evidence-based psychological therapy that has been used to improve...
physical and mental health among adults with long-term conditions, including cancer. It fosters engagement with, rather than avoidance of, painful experiences, to move towards acceptance of unchangeable difficulties alongside building a rich and meaningful life despite the presence of ongoing problems. Furthermore, the acceptance-type responses encouraged in ACT are associated with improved post-treatment functioning compared with other response styles, such as suppression, in adult patients with brain tumour. This gives ACT face validity for application to brain tumour survivors, where there can be permanent cognitive impairment and unavoidable, ongoing physical symptoms. DNA-v is an adaptation of ACT that is showing greater acceptability among young people. However, there has been no published ACT intervention research with young people surviving brain tumours to date.

Many young people who survive brain tumours receive care on a regional basis from principal treatment centres which are often a long journey from home, challenging equity of access to psychological care. Regular psychological therapy sessions at such centres cause additional disruption to patients’ lives during the post-treatment phase, when they are typically aiming to re-engage with normal life. Remote delivery of psychological therapy can improve accessibility to treatment for those who would not otherwise seek psychological care and outcome-effectiveness can be similar to face-to-face treatment. In addition, during the COVID-19 pandemic there have been calls for mental healthcare to be enhanced for vulnerable groups specifically including the use of remotely delivered mental health treatment.

In summary, the psychosocial needs of people surviving childhood brain tumours are often not explicitly treated, resulting in poorer quality of life. A suitable modality for a psychological intervention is ACT, due to its focus on living well despite continuing difficulties. Given the ongoing COVID-19 restrictions and geographical organisation of brain tumour services, remote therapy is likely to be the most suitable and accessible delivery medium.

This study aims to assess the feasibility and acceptability of ACT as a psychological treatment to improve quality of life among adolescents and young adults after brain tumour treatment. Active intervention will be compared with a waitlist control group after 12 weeks follow-up. The durability of treatment effects will be assessed over a 12-month follow-up period. The intervention will be delivered via video conferencing and will therefore also assess the feasibility and acceptability of video conferencing delivery of ACT for this population. The secondary aim will be a preliminary assessment of clinical and cost-effectiveness of ACT.

**METHOD**

**Design**

This study is a two-arm, parallel group, randomised controlled trial comparing ACT with waitlist controls who will then receive ACT after the 3-month waiting period. Participants will be randomised on a 1:1 ratio to receive 12 weeks of treatment either immediately or following a 12-week wait. The treatment will be DNA-v which is a model of ACT adapted to those aged 11–24 years. The model will be further adapted for those who have undergone brain tumour treatment. As trial therapists will perform assessments, blinding is not possible. Follow-up assessments will be conducted at 12 weeks, 24 weeks, 36 weeks and 48 weeks post-randomisation with primary endpoint at 12 weeks. The later follow-ups will be used to assess the durability of any clinical effects. No restrictions will be placed on access to other services during study participation, but service use will be recorded. Recruitment started 04 January 2021 and is planned to end 31 January 2023.

A subsample of participants will be invited to participate in an embedded process evaluation, which will include semi-structured interviews with 7–10 participants from each arm of the trial after their post-treatment follow-up. Purposive sampling will be used to gain representation across trial arms, age, gender, ethnicity, intervention engagement and intervention satisfaction. This qualitative data will be complemented by post-session reflective video or written diary data, in which participants will briefly reflect on each session and what they found most helpful (figures 1 and 2).

**Patient and public involvement**

Young people with lived experience of brain tumours have been involved in the study from the initial funding application and review. Since then, a group of nine young people with lived experience of brain tumours have contributed to a patient advisory group, facilitated by the chief investigator and trial manager and assisted by a member of the brain tumour charity patient support team. All key decisions in developing and running the trial involve separate discussion with the Patient Advisory Group prior to discussion with the trial management group (TMG; which also includes PAG membership). The PAG has opportunity to raise ideas, concerns or questions independent of the TMG agenda.

Specifically, patient feedback has been used to design the recruitment and assessment processes to ensure that there is sufficient opportunity for participants to choose the level of parental involvement, particularly for those aged 16–24. Given the ongoing inclusion of patient representatives in the research team, their advice will inform dissemination plans that will enable the greatest reach to relevant patient groups and services.

**Participants**

Using a reference study of ACT for adolescents, to detect a clinically important average difference of 0.5 SD between intervention and waitlist control with 80% power at p<0.05 (two-tailed), requires a sample size of 72 participants allowing 20% dropout.
Inclusion criteria
Participants will be included if they meet the following criteria:
► Aged 11–24 years at the time of randomisation.
► Received treatment for a brain tumour at a participating principle treatment centre.
► Active brain tumour treatment is complete and their condition stable for at least 6 months
► Have sufficient cognitive ability to engage with ACT sessions as judged by the clinician at baseline assessment.
► Competent to provide informed consent (participants aged 16 or over) or assent (participants aged 11–15).
► Parent/carer competent to provide informed consent (for participants aged 11–15).

Exclusion criteria
Participants will be excluded if affected by the following criteria:
► Received a structured behavioural intervention within 6 months prior to study recruitment.
► Previous or current alcohol/substance dependence, psychosis, suicidality or eating disorder
► Moderate or severe intellectual disability, confirmed through researcher judgement at screening through questions relating to school type and previous diagnoses.
► Immediate risk to self or others.
► The patient or their parent/carer is not able to speak, read or write English.

Figure 1 Trial flowchart. ACT, acceptance and commitment therapy; REDCap, Research Electronic Data Capture.
Study setting and recruitment

Participants will be recruited from one of three usual care oncology teams based at three principle treatment centres covering different regions of England. Oncology staff will approach eligible patients and seek consent for a researcher to contact them. Participants will also be recruited via public advertisements in hospital clinics and national cancer charity publicity. If consent to contact is
given, or patients make contact via advertising, patients (and parents/carers for patients under 16) will be sent age-appropriate study information and a screening assessment will be arranged to assess eligibility. At the screening assessment informed consent will be sought either remotely or face-to-face prior to enrolment (see supplementary materials for model consent forms). Patients will then be randomised at enrolment using the REDCap (Research Electronic Data Capture; https://www.redcapcloud.com/) secure cloud storage system. Researchers completing assessments will be qualified clinical psychologists.

**Assessments**

**Screening**
The screening questionnaire assesses patient eligibility. Where patients are aged over 16 they will be given the choice of whether they wish to involve their parent/carer initially and verified separately with the patient alone. The 5 min screening questionnaire asks about brain tumour history and treatment, contact with healthcare services, other diagnoses/problems, type of education facility attended, age of participant and other eligibility criteria. Patients who do not meet screening criteria because they have recently received a behavioural therapy within the specified exclusion time frame (6 months), may be eligible for rescreening at a later date when these time exclusions have passed. In which case, researchers will seek consent to contact these patients at an agreed later point to reassess eligibility.

**Feasibility**

Feasibility will be assessed by documenting the proportion of patients showing interest who then consent to the trial and complete the intervention. Completion is defined as attending five or more ACT sessions. Fidelity to the ACT therapeutic model will be assessed monthly and then bimonthly using the ACT fidelity measure.

**Acceptability**

Acceptability will be assessed using the session attendance rate. An adapted 6-item patient-reported version of the Credibility/Expectancy Questionnaire will also be given at baseline and after the second ACT session to assess treatment credibility. Participants’ experience of the intervention will be assessed using the Experience of Service Questionnaire, with an additional item to assess video conferencing treatment satisfaction.

**Session-by-session routine outcome monitoring**

Routine outcome monitoring involves monitoring change in therapeutic outcome session-by-session and offering therapists feedback on algorithm-based predicted post-treatment changes using similar outcome trajectories of previous patients. The approach is evidenced as best practice within psychological therapies in general and specifically for children and adolescents, because it reduces therapy failure, increases the speed of improvements, and may enhance the magnitude of improvement.

Each ACT session will begin by completing the Outcome Rating Scale: a brief, general assessment of self-reported well-being. Each session will end by completing the Session Rating Scale: a brief assessment of therapeutic alliance.

**Psychological flexibility**

Psychological flexibility is the main treatment target in ACT, which is the ability to remain in contact with current experiences even when unwanted thoughts, feelings or sensations are present, while selecting actions that support one’s personal values. Psychological flexibility is associated with improved physical and mental health, quality of life and functioning.

**Acceptance and Action Questionnaire II**

The Acceptance and Action Questionnaire II (AAQ-II) is a 7-item self-reported assessment of psychological inflexibility for participants aged 16 and over. (eg, ‘emotions cause problems in my life’).

**Avoidance and Fusion Questionnaire for Youth 8-items**

The Avoidance and Fusion Questionnaire for Youth 8-items (AFQ-Y8) will be used as an 8-item self-reported measure of psychological inflexibility for those aged 11–15 or older participants with mild intellectual disabilities (eg, ‘my thoughts and feelings mess up my life’).

**Mental health and well-being**

**WHO Well-Being Index 5-items**

The WHO Well-Being Index 5-items (WHO-5) is a 5-item self-reported assessment of well-being and mental health suitable for children and adults (eg, ‘over the past week, I have felt calm and relaxed’).

**Generalised Anxiety Disorder assessment 7-items**

The Generalised Anxiety Disorder 7-items (GAD-7) is a 7-item self-reported measure of generalised anxiety symptoms based on diagnostic criteria (eg, worrying too much about different things over the previous 2 weeks).

**Patient Health Questionnaire 9-items**

The Patient Health Questionnaire 9-items (PHQ-9) assesses depression symptoms on nine self-reported items based on diagnostic criteria for major depression (eg, feeling down depressed or hopeless over the previous 2 weeks). The PHQ-9 is validated for use among adolescents and adults.

**General health and quality of life**

**EuroQol 5-dimensions 3-levels**

The EuroQol 5-dimensions 3-levels (EQ-5D-3L) is a self-reported assessment of health-related quality of life for participants aged 16–24. Each dimension is rated at 3-levels: no problems, some problems and extreme problems (eg, ‘I have no problems with self-care’).
The DNA+ therapeutic model is an adaptation of ACT specifically for adolescents and young adults, which aims to be developmentally relevant for this stage of life. The approach involves exploration of different skills to address difficulties. Specifically, the approach teaches participants how to manage their behaviour by grouping them into four skills. First, the V, stands for vitality and participants are coached in the use of video conferencing software to maximise therapeutic benefits. This will include fluent use of screen-sharing and sharing computer software facilities. This aims to support therapists and participants to use all activities that might be included in a face-to-face consultation when working remotely. Education will also involve agreement on simple contingency management plans for managing video conferencing connection failure and agreed signals if confidentiality is compromised.

The EuroQol 5-dimensions youth-version (EQ-5D-Y) is a modified version that combines elements of the child and adolescent service usage schedule, 35 such as school attendance. The CSRI is completed with the parent/carer unless the participant is seeking treatment independently, in which case the participant completes the CSRI.

Routine outcome monitoring is completed at each ACT session and clinical outcomes are collected at baseline, 12-week, 24-week, 36-week and 48-week follow-up for the PHQ-9, GAD-7, WHO-5, AAQ-II/AFQ-Y8, EQ-5D and PROMIS. The CSRI and the SDQ will only be completed at 12-week and 48-week follow-up (see table 1 for the schedule of assessments). All anonymised outcome data will be collected and stored on REDCap. Data management procedures will be aligned with those of the sponsoring NHS Trust.

The DNA+ therapeutic model is an adaptation of ACT specifically for adolescents and young adults, which aims to be developmentally relevant for this stage of life. The approach involves exploration of different skills to address difficulties. Specifically, the approach teaches participants how to manage their behaviour by grouping them into four skills. First, the V, stands for vitality and value and is used to help guide choices toward living with meaning despite difficult experiences such as cancer. ‘The Discoverer’ is used to learn how one can interact with the world on a trial and error basis, investigating and discovering through building strengths and creating new behaviours. ‘The Noticer’ describes one’s observing experiences and learning to respond with awareness rather than being reactive; exploring with senses, body and awareness. ‘The Advisor’ is the skill of learning how we use our inner voice to problem solve, make sense of situations and make decisions based on previous experiences and learning history. Participants are encouraged to move between these roles and investigate what it is like to approach situations, including times of difficulty, in different ways. The overall aim is to learn how to choose actions that support a meaningful, purposeful life.

Involvement of parents/carers will be incorporated within the treatment protocol, particularly for younger participants. The level of parent/carer involvement will be negotiated individually. The treatment is primarily focused on individual work with the participant and joint session time intends to feedback learning to parents/carers with the aim of negotiating ways of integrating helpful aspects of therapy into home life.

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Using online software (https://www.myoutcome-sukapp.com/) throughout therapy, brief measures of patient rated outcome and patient experience of each session will be used to evaluate and guide treatment progress. These processes ensure that therapeutic problems are identified and addressed quickly.

Personalised, targeted smart messaging may improve retention in psychological therapy and outcomes after the completion of therapy. Therefore, relapse prevention plans will be organised into brief weekly reminders that can be sent as text messages after therapy has finished. The messages received will be tailored to participant-reported well-being each week, with messages specific to times when they are doing well, experiencing early warning signs of relapse or experiencing full relapse.

**METHOD OF ANALYSIS**

**Primary outcome analysis**

All baseline outcome analysis will be summarised by randomised group. Categorical data will be reported as frequencies (%) and continuous data will be reported as mean (SD); unless skewed then they will be reported as median (IQR). Any baseline characteristics that are seen to differ

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**Visual Analogue Scale**

The Visual Analogue Scale (VAS) is a self-reported assessment of general health suitable for all participants using a 0–100 VAS, with 0 defined as the worst health imaginable and 100 as the best health imaginable.

**Patient-Reported Outcomes Measurement Information System, Satisfaction with Social Roles and Activities**

The Client Service Receipt Inventory (CSRI) is a flexible research instrument developed to collect information on service receipt, service-related issues and income. This modified version also combines elements of the child and adolescent service usage schedule, 35 such as school attendance. The CSRI is completed with the parent/carer unless the participant is seeking treatment independently, in which case the participant completes the CSRI.

**Health service usage**

The Strengths and Difficulties Questionnaire (SDQ) is a 25-item patient-completed measure of behavioural and emotional functioning (and parent/carer-completed where the patient is under 16).

**Functioning**

The Strengths and Difficulties Questionnaire 25-item is a completed measure of behavioural and emotional functioning (and parent/carer-completed where the patient is under 16).

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**Assessment schedule**

Routine outcome monitoring is completed at each ACT session and clinical outcomes are collected at baseline, 12-week, 24-week, 36-week and 48-week follow-up for the PHQ-9, GAD-7, WHO-5, AAQ-II/AFQ-Y8, EQ-5D and PROMIS. The CSRI and the SDQ will only be completed at 12-week and 48-week follow-up (see table 1 for the schedule of assessments). All anonymised outcome data will be collected and stored on REDCap. Data management procedures will be aligned with those of the sponsoring NHS Trust.

**Treatment procedures**

The DNA+ therapeutic model is an adaptation of ACT specifically for adolescents and young adults, which aims to be developmentally relevant for this stage of life. The approach involves exploration of different skills to address difficulties. Specifically, the approach teaches participants how to manage their behaviour by grouping them into four skills. First, the V, stands for vitality and value and is used to help guide choices toward living with meaning despite difficult experiences such as cancer. ‘The Discoverer’ is used to learn how one can interact with the world on a trial and error basis, investigating and discovering through building strengths and creating new behaviours. ‘The Noticer’ describes one’s observing experiences and learning to respond with awareness rather than being reactive; exploring with senses, body and awareness. ‘The Advisor’ is the skill of learning how we use our inner voice to problem solve, make sense of situations and make decisions based on previous experiences and learning history. Participants are encouraged to move between these roles and investigate what it is like to approach situations, including times of difficulty, in different ways. The overall aim is to learn how to choose actions that support a meaningful, purposeful life.

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**METHOD OF ANALYSIS**

**Primary outcome analysis**

All baseline outcome analysis will be summarised by randomised group. Categorical data will be reported as frequencies (%) and continuous data will be reported as mean (SD); unless skewed then they will be reported as median (IQR). Any baseline characteristics that are seen to differ
between the groups will be included in multivariate models to control for confounding by these variables.

Consent will be obtained to use the data of participants who discontinue up until the point they leave the trial. All analysis will be based on observed data, no missing data will be imputed.

The primary outcomes are summarised descriptively:
1. The percentage of patients eligible and approached for the trial who went on to receive at least one session of treatment.
2. The percentage of patients completing treatment (attending at least five treatment sessions).
3. Patient rated experience of the treatment, benchmarked against national database ratings of National Health Service (NHS) child and adolescent psychological therapy services.

**Secondary outcome analysis**

Assuming normally distributed and balanced data, independent t-tests will compare clinical outcomes between arms at 3-month follow-up as the primary endpoint. If there are imbalances between arms multiple linear regression will be used to control for baseline differences.

To assess whether post-treatment changes are maintained during the follow-up period one-way repeated measures analysis of variances will be completed separately for each arm across 9-month and 6-month post-treatment follow-up for the two arms.

If the baseline characteristics of participants in the two arms are imbalanced and there is significant clustering of covariance within participants across assessment time points multilevel modelling will be used to nest follow-up outcomes within reporting participants hierarchically. This accounts for both within and between participant change over time giving a more accurate estimate of change across follow-up. Where the participant sample may be split by age (eg, the AAQ-II and AFQ-Y8) logistic regression will be used to report the OR of achieving minimal clinically important improvement.

**Table 1** Standard Protocol Items: Recommendations for Interventional Trials diagram of assessments at enrolment, allocation, weekly sessions and follow-up

<table>
<thead>
<tr>
<th>Study period</th>
<th>Time points</th>
<th>Enrolment</th>
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<th>Post-allocation</th>
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<tr>
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<td>ESQ</td>
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AAQ-II, Acceptance and Action Questionnaire-II; ACT, acceptance and commitment therapy; AFQ-Y8, Avoidance and Fusion Questionnaire for Youth; CEQ, Credibility/Expectancy Questionnaire; CSRI, Client Service Receipt Inventory; EQ-5D-3L, EuroQol 5-dimensions 3-levels; EQ-5D-Y, EuroQol 5-dimension youth version; ESQ, Experience of Service Questionnaire; GAD-7, Generalised Anxiety Disorder 7-items; ORS, Outcome Rating Scale; PHQ-9, Patient Health Questionnaire 9-items; PROMIS, Patient-Reported Outcomes Measurement Information System – Satisfaction with Social Roles and Activities; SDQ, Strengths and Difficulties Questionnaire; SRS, Session Rating Scale; ; WHO-5, WHO Well-Being Index 5-item.
improvement between treatment group and waitlist control at 3-month follow-up. Logistic regression analyses will also account for group-level imbalances at baseline.

Qualitative methods
Qualitative analysis will be used on purposively sampled interviews and reflective written or video diary entries made by participants directly after each ACT session. Participants will be invited to either remain on the video call after the therapist has left or type on the reflective diary template brief reflections on the session and what they took from it. This qualitative data will be analysed using thematic analysis. Thematic analysis has been selected as a flexible analytical method which allows for both inductive (emerging from the text) and deductive (guided by theory) coding. Interviews and video diary entries will be recorded and transcribed verbatim. Data will be analysed using NVivo to establish themes and subthemes. A theme template will be established and an independent researcher will be asked to code a selection of text extracts to establish the trustworthiness of themes. The theme template will be revised following the independent data audit and consensus discussion among researchers. A mixed methods approach will be used to integrate qualitative and quantitative data in order to explore full implementation of the intervention. This will include integration of the participants’ evaluation of service questionnaire with interview transcript data, diary reports and session attendance rate. This triangulation aims to use the reciprocal effect of session attendance, evaluation of treatment and qualitative reports to gain a fuller understanding of processes facilitating or impeding the feasibility of the intervention.

Economic evaluation
The primary cost-effectiveness analysis will be the mean incremental cost per quality-adjusted life-year of waitlist control compared with active treatment at 12-week follow-up, from an NHS and personal social services cost perspective. This will be derived from the health utility index using the EQ-5D and service use costs using the CSRI.

To calculate the mean cost per patient of ACT we will collect detailed information on therapist time and evaluate the cost of remote delivery. To establish the cost of usual treatment and other healthcare costs in both the intervention and comparator arms additional health and social care resource use will be obtained from a modified version of the CSRI. Unit costs will be taken from national published sources.

For each resource used we will report descriptive statistics for the percentage number of patients that accessed the service and the mean and SD of times they accessed. As part of the secondary analyses we will include cost of education and out-of-pocket costs.

ETHICS AND DISSEMINATION
East Midlands, Nottingham 1 Research Ethics Committee gave ethical approval (Reference: 20/EM/0237).

Trial monitoring and safety
The TMG consists of the study co-investigators and at least two patient advisors with lived experience of brain tumours. The TMG will meet at least every 6 months to discuss study progress and the trial conduct.

Trial steering committee
The trial steering committee (TSC) is formed of a medical statistician, an oncologist, clinical psychologists and is chaired by a clinical academic neuropsychologist. The TSC will provide independent supervision for the trial on behalf of the trial sponsor and funder. Every 6 months the TSC will review: trial progress, adherence to the protocol, patient safety and the consideration of new information of relevance to the research question.

The trial funders and sponsor will not contribute to the collection of data, study management, analysis or interpretation of data, writing of reports or the decision to submit reports for publication (sponsor email address: researchsponsor@nuh.nhs.uk). Any changes to the study protocol would need to be agreed by the TMG, the TSC and the research ethics committee.

Recording adverse events
This study is deemed to be low risk in terms of adverse events related to treatment. There are no drug treatments involved and the therapy trialled is well-established as safe and effective. Nonetheless, serious adverse events (SAEs) that are unexpected and directly related to the intervention will be recorded in the medical records or other designated place following consent. These events will be recorded with clinical symptoms and accompanied with a description of the event. Safeguarding issues will be recorded and reported to the chief investigator immediately and appropriate local NHS safeguarding procedures will be followed.

Reporting SAEs
Participants will be asked to contact the study site immediately after any adverse events and will be asked if any have occurred at follow-up assessments. Any adverse events reported will be assessed to see if it meets the criteria for an SAE that is unexpected and directly related to the intervention. All treatment-related SAEs will be recorded and reported to the research ethics committee as part of the annual reports. Unexpected SAEs will be reported within stipulated time frames. Participants who experience an SAE that is unexpected and related to the intervention may be withdrawn from the study at the discretion of the investigator and sponsor. Every withdrawal will be examined by the TSC.

Dissemination
The dissemination strategy employs a number of routes with aim of maximising the reach to study stakeholders.
Findings will be reported in peer-reviewed journals and at related academic and clinical conferences. Results will also be disseminated to third sector organisations that offer support to young people who have experienced brain tumours. An end of study dissemination event will be held, which will include patients, carers, researchers, clinicians, clinical service managers and commissioners to support ongoing development beyond feasibility, if appropriate, in ways that are most relevant to patients and health services.

**DISCUSSION**

This study will provide an initial assessment of acceptability and feasibility of remotely delivered ACT as a psychological treatment to support young people after brain tumour treatment, coupled with initial evidence of cost and clinical effectiveness. The study will also obtain rich data on patient experience of remote delivery of psychological therapy, which will offer important insights on patients’ perspectives of new ways of working. Overall, this study will provide sufficient evidence to evaluate whether a larger-scale trial is warranted and will inform future trial design.

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4University Hospitals Bristol and Weston NHS Foundation Trust, Bristol, UK
5Population Health Sciences, University of Bristol, Bristol, UK
6Newcastle upon Tyne Hospitals NHS Foundation Trust, Newcastle upon Tyne, UK
7College of Life and Environmental Sciences, University of Exeter, Exeter, UK
8Faculty of Health and Medical Sciences, University of Surrey, Guildford, UK
9Great Ormond Street Hospital For Children NHS Foundation Trust, London, UK
10Children’s Brain Tumour Research Centre, University of Nottingham, Nottingham, UK
11Newcastle University, Newcastle upon Tyne, UK
12NHS Dumfries and Galloway, Dumfries, UK
13DNA-v International, Melbourne, Victoria, Australia

**Acknowledgements** The authors would like to thank all members of the Brain Tumour Charity’s Young Ambassadors who have contributed to the design and development of this study. The authors thank Dr Andrea Venn, University of Nottingham School of Medicine statistics service, for her support in designing the analysis.

**Contributors** ST, RO, IW, HB, JL, FG, RGG, SCC, SB and SL contributed to the conception, planning, design and grant application for the study. ST, SM, RO, LH and JL contributed to designing the intervention and related processes. All authors contributed drafting, revising and approving the manuscript.

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**Competing interests** None declared.

**Patient consent for publication** Not required.

**Provenance and peer review** Not commissioned; peer reviewed for ethical and funding approval prior to submission.

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**ORCID iD**

Sam Malins http://orcid.org/0000-0001-9570-196X

**REFERENCES**


Participant Consent Form (adult)

Version: 0.3  Date: 27/10/2020

Acceptance and Commitment Therapy for Young Brain Tumour Survivors:
An Acceptability and Feasibility Trial
Principal Investigator: Dr Sophie Thomas

Patient Study ID: ..................  Initials: ..................

1. I confirm that I have read and understand the information sheet dated _________ (version _______) for the above study and have had the opportunity to ask questions.

2. I understand and agree that my participation is voluntary and that I am free to withdraw at any time without my medical care or legal rights being affected.

3. I understand and agree that my medical records may be looked at by authorised individuals from the Sponsor for the study and the UK Regulatory Authority in order to check that the study is being carried out correctly.

4. I understand and agree that if I withdraw from the above study, the data collected from me will be used in analysing the results of the trial.

5. I consent to the storage, including electronic, of my personal information for the purposes of this study. I understand that any information that could identify me will be kept strictly confidential and that no personal information will be included in study reports or other publications.

6. Optional: I agree to my treatment sessions being audio-video recorded for the purposes of supervision and research.

7. Optional: I agree to be approached for an audio-video recorded interview about my experiences of treatment in the study.

8. I understand and agree that recordings may be used by researchers for the purpose of ensuring that the treatment is being delivered appropriately or transcribed for research to enable a better understanding of the treatment.

9. I agree that my GP, or any other treating doctor, will be notified of my participation in this study

10. I agree to take part in the study.

11. Optional: I agree that my research data may be stored for possible use in future research during and after 5 years, and shared with other researchers. Any data used will be anonymous, and I will not be identified.

Name of the participant (Print)  date  Participant’s signature

Name of person receiving consent (Print)  date  Signature

Original to be retained and filed in the site file. 1 copy to patient, 1 copy to be filed in patient's notes
Acceptance and Commitment Therapy for Young Brain Tumour Survivors: An Acceptability and Feasibility Trial

(ACT NOW)

IRAS Number: 266746

R&I reference number: 19CS023

Verbal Consent Script

<table>
<thead>
<tr>
<th>Participant ID:</th>
<th>Initials:</th>
</tr>
</thead>
<tbody>
<tr>
<td>Date of Birth:</td>
<td>NHS/Hospital Number:</td>
</tr>
<tr>
<td>Principal Investigator:</td>
<td></td>
</tr>
<tr>
<td>Date of verbal consent call:</td>
<td><strong>/</strong>/____ (dd/mmm/yyyy)</td>
</tr>
<tr>
<td>Time of verbal consent call:</td>
<td><strong>/</strong> (00.00 hr)</td>
</tr>
<tr>
<td>Format of verbal consent e.g. Skype, phone call:</td>
<td></td>
</tr>
</tbody>
</table>

Telephone/contact the participant and confirm that you are speaking to the relevant person, and that it is a convenient time to call (if not, establish if the participant wishes to be contacted at an alternative time; or whether the participant no longer wishes to be contacted regarding the study). Then continue with the script below:

Hello [name of participant], my name is [name of member of staff taking consent] and I am a [role of member of staff e.g. research clinical psychologist] working on the ACT NOW study, which you [have shown interest in / have been given information about / are currently taking part in].

You may have received some documents from us recently regarding the ACT NOW research study and we wondered if you have had a chance to read them? (Allow participants to respond – if they have read it and no longer wish to be contacted with regards to this study, then thank them for their time and say goodbye – if they are unsure or more positive then continue below).

I am calling today to invite you to take part in a treatment study. We would like to see if a talking treatment called Acceptance and Commitment Therapy helps young people who have had a brain tumour. To do this we would like to...
offer you up to 12 one hour sessions of Acceptance and Commitment Therapy and ask you to complete some questionnaires four times over a year.

The Ethics Committee, whose role it is to scrutinise research and protect patients, has agreed that we can obtain verbal consent from you over the phone, but please let me reassure you that whatever you decide, it will not change your treatment in anyway. Do you have any questions you would like to ask me at this stage?

Record participant’s response: Yes / No

If yes, record any questions and responses given, below:

________________________________________________________________________
________________________________________________________________________
________________________________________________________________________
________________________________________________________________________
________________________________________________________________________

Just so that I can check that I have explained myself clearly to you, can you please confirm if you understand what I have told you?

Record participant’s response: Yes / No

If you are happy with my responses, can you please let me know whether you agree or not, to take part in the ACT NOW study?

Record participant’s response: Yes / No

(If participant answered yes, continue overleaf; if no, then thank the participant for their time and say goodbye)

Finally, could I please ask you to confirm a few details to record your consent?

Version and date of information sheet received by participant: Version: _____ Dated: ___ / ____ / _____

Date the study information was received (dd/mmm/yyyy)? ___ / ____ / ____

We would like to send you a copy of the consent form to keep for your information. Could you please confirm for me how you would like to receive this, via post or email? Please confirm your contact details for me:

Address: ..........................................................................................
..........................................................................................
..........................................................................................

Email: ..........................................................................................

ACT NOW
IRAS Number: 266746
act_now_protocol supplementary materials 210520
Page 3 of 16
## PARTICIPANT VERBAL CONSENT CONFIRMATION FORM

**IRAS Number:** 266746  
**R&I reference number:** 19CS023  
**Please initial (researcher):**

<table>
<thead>
<tr>
<th></th>
<th>Yes</th>
<th>No</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. After explaining the study information to the participant, I can confirm that the participant understands and agrees to participate in the above research study.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>2. The participant understands and agrees that the study is voluntary, and that they can withdraw at any time without their medical care or legal rights being affected; although data and samples already collected will be retained for use in the study.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>3. The participant understands and agrees that their medical records may be looked at by authorised individuals from the Sponsor for the study and the UK Regulatory Authority in order to check that the study is being carried out correctly.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>4. The participant understands and agrees that if they withdraw from the study, the data collected from them will be used in analysing the results of the trial.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>5. The participant understands and agrees to the storage, including electronic, of their personal information for the purposes of this study. They understand that any information that could identify them will be kept strictly confidential and that no personal information will be included in study reports or other publications.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>6. <strong>Optional:</strong> The participant agrees to their treatment sessions being audio-video recorded for the purposes of supervision and research.</td>
<td></td>
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<td>7. <strong>Optional:</strong> The participant agrees to be approached for an audio-video recorded interview about their experiences of treatment in the study.</td>
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<td>8. The participant understands and agrees that recordings may be used by researchers for the purpose of ensuring that the treatment is being delivered appropriately or transcribed for research to enable a better understanding of the treatment.</td>
<td></td>
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<tr>
<td>9. The participant understands that their GP or any other treating doctor, may be notified about their participation in the study.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>10. <strong>Optional:</strong> The participant understands and agrees that their research data may be stored for possible use in future research during and after 5 years, and shared with other researchers. Any data used will be anonymous, and they will not be identified.</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
Participant

Full name *(block capitals)*: 

Date/time verbal consent provided by participant: 

<table>
<thead>
<tr>
<th>DD/MMM/YYYY</th>
<th>Time (24 hr)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Person taking consent

I have explained the study to the above named participant and they have indicated their willingness to participate.

Full name *(block capitals)*: 

Signature: 

Date: 

<table>
<thead>
<tr>
<th>DD/MMM/YYYY</th>
</tr>
</thead>
<tbody>
<tr>
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</tbody>
</table>

Original to be retained and filed in the site file, one copy for the participant, and one copy to be filed in the participant’s medical notes.
Participant Consent Form (Parents/Careers)

Version: 0.3 Date: 27/10/2020

Patient Study ID: ………………… Initials: …………………

Patient initial each box

1. I confirm that I have read and understand the information sheet dated ____________ (version ______) for the above study and have had the opportunity to ask questions.

2. I understand and agree that my child’s participation is voluntary and that they are free to withdraw at any time without their medical care or legal rights being affected.

3. I understand and agree that my child’s medical records may be looked at by authorised individuals from the Sponsor for the study and the UK Regulatory Authority in order to check that the study is being carried out correctly.

4. I understand and agree that if my child withdraws from the above study, the data collected from them will be used in analysing the results of the trial.

5. I consent to the storage, including electronic, of my child’s personal information for the purposes of this study. I understand that any information that could identify my child will be kept strictly confidential and that no personal information will be included in study reports or other publications.

6. **Optional:** I agree to my child’s treatment sessions being audio-video recorded for the purposes of supervision and research.

7. **Optional:** I agree to my child being approached for an audio-video recorded interview about their experiences of treatment in the study.

8. I understand and agree that recordings may be used by researchers for the purpose of ensuring that the treatment is being delivered appropriately or transcribed for research to enable a better understanding of the treatment.

9. I agree that my child’s GP, or any other treating doctor, will be notified of their participation in this study.

10. I agree for my child to take part in the study.

11. **Optional:** I agree that my child’s research data may be stored for possible use in future research during and after 5 years, and shared with other researchers. Any data used will be anonymous, and my child will not be identified.

Name of the participant’s parent/carer (Print) date Parent/carer’s signature

Name of person receiving consent (Print) date Signature

Original to be retained and filed in the site file. 1 copy to patient, 1 copy to be filed in patient’s notes
## Acceptance and Commitment Therapy for Young Brain Tumour Survivors: An Acceptability and Feasibility Trial (ACT NOW)

**IRAS Number: 266746**

**R&I reference number: 19CS023**

**Verbal Consent Script**

<table>
<thead>
<tr>
<th>Participant ID:</th>
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</tr>
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<tbody>
<tr>
<td>Date of Birth:</td>
<td>NHS/Hospital Number:</td>
</tr>
<tr>
<td>Principal Investigator:</td>
<td></td>
</tr>
</tbody>
</table>

**Date of verbal consent call:** ___/___/____  (dd/mmm/yyyy)

**Time of verbal consent call:** ___/___  (00.00 hr)

**Format of verbal consent e.g. Skype, phone call:**

*Telephone/contact the participant’s parent/carer and confirm that you are speaking to the relevant person, and that it is a convenient time to call (if not, establish if the parent/carer wishes to be contacted at an alternative time; or whether the parent/carer no longer wishes to be contacted regarding the study). Then continue with the script below:*

Hello [name of parent/carer], my name is [name of member of staff taking consent] and I am a [role of member of staff e.g. research clinical psychologist] working on the ACT NOW study, which you and your child [have shown interest in / have been given information about / are currently taking part in].

You may have received some documents from us recently regarding the ACT NOW research study and we wondered if you have had a chance to read them? *(Allow participants to respond – if they have read it and no longer wish to be contacted with regards to this study, then thank them for their time and say goodbye – if they are unsure or more positive then continue below).*
I am calling today to invite your child to take part in a treatment study. We would like to see if a talking treatment called Acceptance and Commitment Therapy helps young people who have had a brain tumour. To do this we would like to offer your child up to 12 one-hour sessions of Acceptance and Commitment Therapy and ask you and your child to complete some questionnaires four times over a year.

The Ethics Committee, whose role it is to scrutinise research and protect patients, has agreed that we can obtain verbal consent from you over the phone, but please let me reassure you that whatever you decide, it will not change your child’s treatment in anyway. Do you have any questions you would like to ask me at this stage?

Record parent/carer’s response: Yes / No

If yes, record any questions and responses given, below:

________________________________________________________________________________________________________________________________________________________________________________________

________________________________________________________________________________________________________________________________________________________________________________________

________________________________________________________________________________________________________________________________________________________________________________________

________________________________________________________________________________________________________________________________________________________________________________________

Just so that I can check that I have explained myself clearly to you, can you please confirm if you understand what I have told you?

Record parent/carer’s response: Yes / No

If you are happy with my responses, can you please let me know whether you agree or not, to take part in the ACT NOW study?

Record parent/carer’s response: Yes / No

(If parent/carer answered yes, continue overleaf; if no, then thank the participant for their time and say goodbye)

Finally, could I please ask you to confirm a few details to record your consent?

Version and date of information sheet received by parent/carer: Version: ___ Dated: ___ / ___ / _____

Date the study information was received (dd/mmm/yyyy)? ___ / ___ / _____

We would like to send you a copy of the consent form to keep for your information. Could you please confirm for me how you would like to receive this, via post or email? Please confirm your contact details for me:

Address: ........................................................................................................................................

........................................................................................................................................

........................................................................................................................................

Email: ........................................................................................................................................
PARTICIPANT VERBAL CONSENT CONFIRMATION FORM (PARENT)
IRAS Number: 266746
R&I reference number: 19CS023

Please initial (researcher)

1. After explaining the study information to the participant’s parent/carer, I can confirm that the parent/carer understands and agrees to their child participating in the above research study.

2. The parent/carer understands and agrees that the study is voluntary, and that their child can withdraw at any time without their medical care or legal rights being affected; although data and samples already collected will be retained for use in the study.

3. The parent/carer understands and agrees that their child’s medical records may be looked at by authorised individuals from the Sponsor for the study and the UK Regulatory Authority in order to check that the study is being carried out correctly.

4. The parent/carer understands and agrees that if their child withdraws from the study, the data collected from them will be used in analysing the results of the trial.

5. The parent/carer understands and agrees to the storage, including electronic, of their child’s personal information for the purposes of this study. They understand that any information that could identify their child will be kept strictly confidential and that no personal information will be included in study reports or other publications.

6. **Optional:** The parent/carer agrees to their child’s treatment sessions being audio-video recorded for the purposes of supervision and research.

7. **Optional:** The parent/carer agrees to their child being approached for an audio-video recorded interview about their experiences of treatment in the study.

8. The parent/carer understands and agrees that recordings may be used by researchers for the purpose of ensuring that the treatment is being delivered appropriately or transcribed for research to enable a better understanding of the treatment.

9. The parent/carer understands that their child’s GP or any other treating doctor, may be notified about their participation in the study.

10. **Optional:** The parent/carer agrees that their child’s research data may be stored for possible use in future research during and after 5 years, and shared with other researchers. Any data used will be anonymous, and they will not be identified.
Participant’s parent/carer

Full name (block capitals):

Date/time verbal consent provided by participant:

Person taking consent

I have explained the study to the above named participant’s parent/carer and they have indicated their willingness for their child to participate.

Full name (block capitals):

Signature:

Date:

Original to be retained and filed in the site file, one copy for the participant, and one copy to be filed in the participant’s medical notes.
## Participant Assent Form

**Acceptance and Commitment Therapy for Young Brain Tumour Survivors: An Acceptability and Feasibility Trial**  
**IRAS Reference: 266746**

**Participant Study ID:** ……………………………………  
**Initials:** ………………………………………

<table>
<thead>
<tr>
<th>Question</th>
<th>Yes / No</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 Has somebody explained the study to you?</td>
<td></td>
</tr>
<tr>
<td>2 Do you understand that it is okay to stop taking part at any time? This will not change the way your doctors treat you.</td>
<td>Yes / No</td>
</tr>
<tr>
<td>3 Do you understand and agree that your medical records may be checked to make sure the study is being done properly?</td>
<td>Yes / No</td>
</tr>
<tr>
<td>4 Do you understand and agree that if you stop taking part the study team will use your information in their results?</td>
<td>Yes / No</td>
</tr>
<tr>
<td>5 Do you understand and agree that the research team will store your personal information for the study? No personal information will be shared in public.</td>
<td>Yes / No</td>
</tr>
<tr>
<td>6 Optional: Do you agree to your treatment sessions being audio-video recorded for the study?</td>
<td>Yes / No</td>
</tr>
<tr>
<td>7 Optional: Do you agree to be approached for an audio-video recorded interview about your treatment in the study?</td>
<td>Yes / No</td>
</tr>
<tr>
<td>8 Do you understand and agree that recordings will be used by researchers to make sure the treatment is done properly and to understand how the treatment works?</td>
<td>Yes / No</td>
</tr>
<tr>
<td>9 Do you understand that the doctors who are treating you will be told that you are taking part in this study?</td>
<td>Yes / No</td>
</tr>
<tr>
<td>10 Are you happy to take part in this study?</td>
<td>Yes / No</td>
</tr>
<tr>
<td>11 Optional: Do you agree to your research information being kept for future research? You will not be identified in any research information</td>
<td>Yes / No</td>
</tr>
<tr>
<td>Your full name:</td>
<td>Signature:</td>
</tr>
<tr>
<td>--------------------</td>
<td>------------</td>
</tr>
<tr>
<td>Name of person with parental responsibility for the patient:</td>
<td>Signature:</td>
</tr>
<tr>
<td>Name of researcher taking consent:</td>
<td>Signature:</td>
</tr>
</tbody>
</table>

When completed: 1 for participant, 1 (original) for Investigator's study file and 1 to be kept in participant's medical notes.
Acceptance and Commitment Therapy for Young Brain Tumour Survivors: An Acceptability and Feasibility Trial

IRAS Number: 266746

R&I reference number: 19CS023

Verbal Consent Script

<table>
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</table>

Date of verbal consent call: ___/___/____ (dd/mmm/yyyy)

Time of verbal consent call: ___/___ (00.00 hr)

Format of verbal consent e.g. Skype, phone call:

Telephone/contact the participant and confirm that you are speaking to the relevant person, and that it is a convenient time to call (if not, establish if the participant wishes to be contacted at an alternative time; or whether the participant no longer wishes to be contacted regarding the study). Then continue with the script below:

Hello [name of participant], my name is [name of member of staff taking consent] and I am a [role of member of staff e.g. research nurse] working on the ACT NOW study, do you remember someone talking to you about the study? (Allow participants to respond, if they do not remember briefly summarise the study, as below).

Have you been sent some information from us about the ACT NOW research study and have you had a chance to read them? (Allow participants to respond – if they have read it and no longer wish to be contacted with regards to this study, then thank them for their time and say goodbye – if they are unsure or more positive then continue below).
I am calling today to invite you to take part in ACT NOW. ACT NOW is a research study where you will be offered a talking therapy to see how it can help young people who have had a brain tumour. A talking therapy is a way of helping people without using medicines. Taking part in the study would mean going to up to 12 one hour sessions of talking therapy over a video-call and completing some questionnaires four times over a year.

The Ethics Committee makes sure we do the research safely. They have agreed that we can ask you if you want to take part over the phone, but you do not have to take part and your treatment will not change no matter what you decide to do. You can also leave the study after you have started if you change your mind. Do you have any questions you would like to ask me?

Record participant’s response: Yes / No

If yes, record any questions and responses given, below:

________________________________________________________________________
________________________________________________________________________
________________________________________________________________________

Just so that I can check that you know what the study involves please could you explain what you understood about the ACT NOW study?

Does the participant’s response confer understanding of the study? Yes / No

If you are happy with what I’ve said, can you please let me know whether you agree or not, to take part in the ACT NOW study?

Record participant’s response: Yes / No

(If participant answered yes, continue overleaf; if no, then thank the participant for their time and say goodbye)

Finally, please can I ask you to confirm a few details to record your consent?

Version and date of information sheet received by participant: Version: ___ Dated: ___ / ____ / _____

Date the study information was received (dd/mmm/yyyy)? ___ / ____ / ____

We would like me to send you a copy of the consent form for you to keep. How would you like to receive this, by post or email? Please confirm your contact details:

Address: ..........................................................................................
..........................................................................................

Email: ..........................................................................................
### PARTICIPANT VERBAL CONSENT CONFIRMATION FORM

**IRAS Number: 266746**  
**R&I reference number: 19CS023**

<table>
<thead>
<tr>
<th>Please Initial (researcher)</th>
<th>Yes</th>
<th>No</th>
</tr>
</thead>
<tbody>
<tr>
<td>11. After explaining the study information to the participant, I can confirm that the participant understands what the study is about and agrees to participate in the above research study.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>12. The participant understands that it is okay to stop taking part at any time and that this will not change the way their doctors treat them.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>13. The participant understands and agrees that their medical records may be checked to make sure the study is being done properly.</td>
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<td></td>
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<td></td>
<td></td>
</tr>
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<td>15. The participant understands and agrees that the research team will store their personal information for the study and that no personal information will be shared in public.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>16. <strong>Optional</strong>: The participant agrees to their treatment sessions being audio-video recorded for the study.</td>
<td>Yes</td>
<td>No</td>
</tr>
<tr>
<td>17. <strong>Optional</strong>: The participant agrees to be approached for an audio-video recorded interview about their treatment in the study.</td>
<td>Yes</td>
<td>No</td>
</tr>
<tr>
<td>18. The participant understands and agrees that recordings will be used by researchers to make sure the treatment is done properly and to understand how the treatment works.</td>
<td>Yes</td>
<td>No</td>
</tr>
<tr>
<td>19. The participant understands that the doctors who are treating them will be told that they are taking part in the study.</td>
<td>Yes</td>
<td>No</td>
</tr>
<tr>
<td>20. <strong>Optional</strong>: The participant agrees to their research information being kept for future research and that they will not be identified in any research information.</td>
<td>Yes</td>
<td>No</td>
</tr>
</tbody>
</table>
Nottingham University Hospitals NHS

Participant

Full name (block capitals):

Date of Birth:

Date/time verbal consent provided by participant:

Person taking consent

I have explained the study to the above named participant and they have indicated their willingness to participate.

Full name (block capitals):

Signature:

Date:

Original to be retained and filed in the site file, one copy for the participant, and one copy to be filed in the participant’s medical notes.