Cumbria, Northumberland,
Tyne and Wear

[Insert local site logos here]
IRAS Number: 323903 Study Title: MUSE ARMS Feasibility Trial.
Parent/Guardian Consent Form [Version 2.0 23022023]

Centre Name: [e.g. CNTW / Other NHS Participating Organisation]

Participant ID Number:

MUSE ARMS Feasibility Trial PARENT/GUARDIAN INFORMED CONSENT FORM

	Initial b		
	†	to agree	
2	I confirm that I have read the parent/guardian information sheet dated		
	affected.		
3	I understand that if my child withdraws from the study, or needs to be withdrawn due to becoming too unwell, the research team will keep the research data about my child that they already have, and if you give consent to question 10 on this form they will continue to track long term outcomes via the MHSDS/medical notes unless you request that they do not.		
4	I understand that relevant sections of my child's medical notes and data collected during the study, may be looked at by responsible individuals from [Research site] and from the research Sponsor CNTW NHS Foundation Trust, their representatives and regulatory authorities for the purposes of this research study, which includes audit and monitoring for research quality assurance. I give permission for these individuals to have access to these records in accordance with this study participant information sheet and informed consent and my child's agreement (assent).		
5	I understand and agree that the information collected about my child in the course of this study will be held and maintained by [enter name of organisation(s) that will be storing the participant data], and CNTW and archived at [enter name of organisation(s)] and CNTW.		
6	I agree to my child's NHS Care Team being informed of their participation in the study.		
7	I agree for a brief summary of the research assessments and any treatment sessions to be shared with my child's clinical team (i.e. added into NHS care notes).		
8	OPTIONAL: I consent to the use of audio recording of my child's treatment sessions, so long as my child agrees to this, to check the quality of the MUSE treatment. I understand recordings will follow NHS data security standards for storage and will be destroyed once they are checked for treatment quality.		
	Circle decision: YES / NO		

Print double sided. When completed: One for participant, one for Investigator Site File, and one to be kept in medical notes.

9	OPTIONAL: I consent to my child to take part in an interview about the trial, if they wish to do this.	heir experience in		
	I recognise not everyone is asked to do this and that my child can change their mind			
	any time. I am aware that these reflective interviews are audio recorded anonymously			
	(using an ID code as identifier) and then transcribed during which any further potential			
	personal identifying information is removed ahead of analysis of res	=		
		ecision: YES / NO		
10	,			
	medical databases to look at long term outcomes including use of he	•		
	services, so long as they are in agreement with this. Medical databa Records, and the Mental Health Services Data Set (MHSDS). This r	•		
	my child's consent form and NHS record number to be sent securely			
	processing and storage in the trial master file, which I agree to.			
11	, , , , , , , , , , , , , , , , , , , ,			
	writing up of research findings. The data in an anonymised format w			
	researchers for this study who have a role in analysing and writing u	p data.		
12	I understand that in accordance with openness of data findings the a	nonymised data		
	set from the study may be published in open access and or for wide	r research. My		
	child's personal details will not be shared.			
13	Lagrage for my shild to take part in the above study if they wish to do			
13	I agree for my child to take part in the above study if they wish to do so.			
14	OPTIONAL: I agree for my child to be contacted with end of study information on the			
	trial and their preferred contact method is: email / post / text message <i>(circle as</i>			
	appropriate). Contact details will be obtained from medical records.			
	Circle de	ecision: YES / NO		
Name o	f Parent/Guardian			
Signature of Parent/Guardian Date				
*I certif	y that the information provided was discussed in a language accessible	e to the Parent/Guard	ian. That	
they retained and understood the information for a sufficient period in order to weigh up their decision and				
commu	inicate their decision regarding informed consent.			
*Name	of Researcher Obtaining Consent			
*Signat	ure of Researcher Obtaining Consent	Date		
	•			

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