(Form to be on headed paper)

IRAS ID:

Centre Number:

Low-dose atropine eye drops to reduce progression of myopia in children: a multicentre placebo controlled randomised trial in the United Kingdom (CHAMP UK)

PARENT INFORMED CONSENT FORM

Chi	ld's name:	-
		Please Initial box
1.	I confirm that I have read and understand the Parent Information Sheet	
	for the above study dated XXXX XXX 20XX (Version X.0) and website	
	dated XX ^{XX} XXX 20XX (Version 1.0) and have been given a copy to	
	keep. I have had the opportunity to ask questions and discuss what the	
	study would involve for my child and understand why the research is	
	being done and any foreseeable risks involved.	
2.	I confirm that I have parental responsibility for the child named above.	
3.	My child has indicated that they would like to participate in this research	
	study and has signed an informed assent form.	
4.	I understand that my child's participation is voluntary and that he/she is	
	free to withdraw at any time, without giving any reason, without his/her	
	eye care or legal rights being affected.	
5.	I understand that my child's medical notes and other patient information	
	systems, and the data collected during the study may be reviewed by	
	responsible individuals from the clinical research facility, local Health	
	and Social Care Trust, trial co-ordinating centre, sponsor or regulatory	
	authorities, where it is relevant to my child taking part in this research. I	
	give permission for these individuals to have access to my child's eye	

	health records. I agree to information retained at the Northern Ireland Belfast Health & Social Care Trus	d Clinical Trials Unit	•			
6.	I have to the best of my knowled child's previous or current illness currently participating in any other	ses and medication.	My child is not			
7.	I give permission for my child's in strict confidence by responsible p		•			
8.	I give permission for my child's eye health checks to be carried out by doctors and optometrists from the study team.					
9.	I agree to the data we provide being used in an anonymised format in publications and at conferences and understand that my child will not be personally identified.					
10. I agree to my child's GP being informed of his/her participation in the study.						
11. I agree to complete some questionnaires with my child five years after the study has ended.						
ag	ree to my child taking part in the	e above study				
lan	ne of Parent	Date	Signature			
lan	ne of Person taking consent	 Date	Signature			