



Remote Participant Consent Form

The POPPY Study - Qualitative study

Patient reported Outcomes, Postoperative pain and Pain relief after daY case surgery

Chief Investigator:

Participant Identification Number for study:

				Person taking consent to <u>initial</u> box	
1.	I confirm that I have read the inform above study. I have had the opport have had these answered satisfacto	tunity to consider the informa			
2.	I understand that my participation is without giving any reason, without r				
3.	. I understand that relevant sections of my medical notes and data collected during the study, may be looked at by individuals from University Hospitals Plymouth NHS Trust, University of Plymouth, from regulatory authorities or from the NHS Trust, where it is relevant to my taking part in this research. I give permission for these individuals, who are bound by a strict duty of confidentiality, to have access to my records.				
4.	I understand that the interview will take place online <i>via</i> Zoom, and I agree for this to be recorded.				
5.	I understand that the information collected about me will be used to support other research in the future, and may be shared anonymously with other researchers.			1	
6.	I understand that the information collected in this study will be presented at scientific meetings and/or published in medical journals. I understand that my identity will not be revealed in any of these cases and my data will be in an anonymous format to protect my identity.				
7.	I agree to take part in the above stu	udy.			
I (name of person taking consent) have read this consent form to (participant name in full) on (date). I confirm that they have:					
	 Understood the participant information sheet and informed consent form and Verbally agreed to the statements numbered 1 to 7. 				
Na	me of person taking consent Da	ate	Signature		

Time of consent (24hr clock): :

When completed: 1 copy for participant (via email/post); original copy for researcher site file.