



Contact : Mr P Gadd
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<i>Study Code:</i>	<i>Site ID Code:</i>	<i>Participant identification number:</i>
<input type="text"/>	<input type="text"/>	<input type="text"/>

PALM CONSENT FORM

Research title: Parkinson's speech to text training through artificial learning methods

Name of Researcher: Paul Gadd

If you agree, please initial box

1. I confirm that I have read the Palm Participant Information Sheet dated 17th Nov 2022 (version 1.3) for this study. I have had the opportunity to consider the information, ask questions and have had these answered satisfactorily.	
2. I understand that my participation is voluntary and that I am free to withdraw at any time without giving any reason, without my medical care or legal rights being affected.	
3. I understand that data collected during the study may be looked at by individuals from Protocol IT Ltd. I give permission for these individuals to have access to my records.	
4. <i>I agree to provide samples of my speech as part of my involvement in this study and I understand I will not gain any direct personal or financial benefit from them.</i>	
5. <i>I agree to audio (speech) recordings being used with a neural network to assist with speech to text recognition and the use of my anonymised feedback in research reports and publications.</i>	
6. <i>I understand that the contact information held and maintained by Protocol IT Ltd may be used to help contact me or provide information about this or future research in the same area.</i>	
7. I agree to take part in this study.	
8. <i>I agree to be contacted about ethically approved research studies for which I may be suitable. I understand that agreeing to be contacted does not oblige me to participate in any further studies.</i>	

<Document title> : PALM Information Consent form	Version/Date: < 1.3 >
<Study Title> Parkinson's speech to text training through artificial learning methods	Project number: < PIT1/2022 >
<Chief Investigator> Paul Gadd	Page: 1 of 2



9. I agree for my anonymised samples to be used in future ethically-approved research in the UK, which may have commercial use.	
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<i>Name of Participant</i>	<i>Date</i>	<i>Signature</i>
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<i>Name of Person taking Consent</i>	<i>Date</i>	<i>Signature</i>
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*1 copy for participant; 1 copy for researcher site file.