CONSENT FORM

Study title		Lessening the impact of fatigue in inflammatory rheumatic diseases: a ran clinical trial (LIFT) IRAS 216267	domised
Name of CI:		Prof Gary J Macfarlane	
Participant ID:			
Plea	Please initial each box		
1.	had the opportunity to o	ead the information leaflet (version, date). I have consider the information, ask questions and have had these answered satisfactorily. In contact the study team if I have any further questions.	
2.	reason, without my mee	participation is voluntary and that I am free to withdraw at any time, without giving any dical care or legal rights being affected. Nation collected prior to my withdrawal will be used in the analysis; however, no new acted.	
3.	individuals from the Uni or from the NHS Trust,	ections of my medical notes and data collected during the study, may be looked at by versity of Aberdeen and collaborators involved in the study, from regulatory authorities where it is relevant to my taking part in this research. hese individuals to have access to my records.	
4.	I agree that identifiable	I by the study team via post, telephone or email after the trial period is over. contact information will be kept after the end of this study and this information will be securely in accordance with the data protection act.	
5.		he LIFT study research group to contact my GP or other health care professional to ng part in this research.	
6.	I give permission for the findings are discovered	he LIFT study research group to notify my GP and/or my health care team if incidental l.	
Please initial one box only			
7.		e of the groups that will get therapist-delivered treatment, I do give permission to the up to record some of the sessions (phone or internet-based audio/video calls) for oses only.	
		of the groups that will get therapist-delivered treatment, I do not give permission to up to record any of the sessions (phone or internet-based audio/video calls).	
Plea	se initial box		Initial
8.	I agree to take part in the second	his randomised study.	
9.	-	nat data collected as part of this study may be linked to with health research datasets search studies. My unique NHS number may be used to do this.	
Please initial one box only			Initial
10. I agree to give additional blood samples for future ethically approved studies, together with the related data, and I understand it may be linked with health research datasets and data from other research studies. My unique NHS number may be used to do this.			
	-	an additional blood sample for future ethically approved studies.	
Name	e of participant	Date (DD/MON/YYYY) Signature	

Name of person taking consent

Date (DD/MON/YYYY)

Signature

NOTE: Original for local Investigator Site File, copies for participant, for central Trial Master File and for participant's medical notes