A multicentre non-blinded randomised controlled trial to assess the impact of \underline{R} egular \underline{E} arly \underline{SPE} cialist symptom \underline{C} ontrol \underline{T} reatment on quality of life in malignant \underline{Meso} the lioma - "RESPECT-Meso"

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Centre Number:	Study Carer Identification	number:	
CARER CONSENT FORM			
	randomised controlled trial to ty of life in malignant <u>Meso</u> th	assess the impact of <u>R</u> egular <u>E</u> elioma - "RESPECT-Meso"	arly <u>SPE</u> cialist symptom
Name of Principal Investigat	or:	<u> </u>	Please initial each box
) dated/_	/ for the above stu	Carer Information Sheet (version dy. I have had the opportunity have these answered satisfactorily	to
	any time without giving any reason, without my medical care or legal rights being		
the study may be authorised by the S Trust, where it is re	looked at by responsible iponsor, regularity authorities,	Il notes and data collected during a collected during the Sponsor the study team or from the NI his research. I give permission for the study team or from the NI his research.	or HS
4. I agree to my GP bei	4. I agree to my GP being informed of my participation in the study.		
5. I agree to take part i	n the above study.		
Name of Carer	 Date	Signature	
Name of person taking Consent	 Date	Signature	

When completed: 1 copy for carer; 1 for study file; 1 (original) for medical notes