Sub study 1 samples for future research - Subject enrollment and identification Log

18.3

Sitenumber: **Site Name:** **Principal Investigator:**

Participants who have signed the additional informed consent for use of excess sub study 1 samples for future research

PID Number (REDCap Number)	**CPR Number**	**Date of signature**	**Patient Name**	**Contact Information (phone) (Only if patient accepts)**

End of trial – *by my signature I certify that the above details are correct*

Signature Principal Investigator	Initials	Date