Request to Conduct Research for Same Use of Stored Human Samples, Specimens, or Data Collected in a Terminated NCI-IRB Protocol

Protocol Number of Terminated Protocol:

Please Check One:

[ ] Exempt from IRB Review:

If the NIH investigator CANNOT identify the subjects, then the research activity may be exempt from IRB review and approval (Under 45CFR46.101(b). Use form I on this OHSR Website http://ohsr.od.nih.gov/info/info.html .

[ ] Same Use as Described in the Terminated Protocol

For this submission, complete the 1195 for as with any other initial protocol, respond to the required new protocol information, and provide a copy of the Consent Form/s from the Terminated Protocol:

I. Brief Description of Data/Specimens (how many, types, storage):

II. Research/Analysis being Conducted:

III. Timeframe for Analysis of Data/Specimens: (Please note – if longer than one year the requirement for Continuing Review of the Research applies.)

IV. Indicate how the Rights and Welfare of Human Subjects will be protected and Risks to Human Subjects will be minimized: (This section must include a description of the processes for maintaining confidentiality of identifiable data/specimens/samples and what will happen to identifiable data/specimens/samples at the end or completion of the protocol. Where human samples or specimens are involved - describe how they will be stored; how they will be tracked; and under what circumstances would the PI report to the IRB loss or destruction of the samples/specimens.