Two circumstances when IRB approved active protocols may be closed by the PI and/or the IRB

The University of Kentucky (UK) Institutional Review Board (IRB) has adopted the following two circumstances permitting IRB approved protocols to be closed by the Principal Investigator (PI) and/or the IRB.

1. **If a study is open and the only activity remaining is analysis of data collected during the study, the study may be closed by the PI if one of the following conditions exist:**
   - the data is de-identified and there are no subject identifying codes or links to the de-identified data; or
   - the data for studies approved after January 21, 2019 with identifiers, is encrypted; or
   - for FDA-regulated studies, there are no outstanding data queries or other investigator/site responsibilities in the trial

2. **If a study has been open for a period of three or more years and there have been no subjects enrolled in the study, the IRB requires that the study be closed.**

   If at Continuation or Annual Administrative Review the investigator reports to the IRB that the study has been open for a period of three or more years and there have been no subjects enrolled into the study, the IRB will require study closure, unless justified (e.g., study involving a rare condition). A notification of closure is sent to the investigator.

The PI must complete and submit a final review report form with the requested materials to the IRB unless:

1. He/she never initiated the study; or,
2. A review (initial, administrative, or continuation) has been conducted within the last six months and no subjects have been enrolled since the last review.

A copy of this handout may be downloaded from the ORI Survival Handbook webpage under the sections Study Closure and Recordkeeping [https://www.research.uky.edu/office-research-integrity/irb-survival-handbook](https://www.research.uky.edu/office-research-integrity/irb-survival-handbook). For additional details on additional study closure circumstances, see the IRB/ORI “Study Closure” Standard Operating Procedure (SOP) available on ORI’s SOP web page: [http://www.research.uky.edu/ori/human/SOPs & Policies.htm#4](http://www.research.uky.edu/ori/human/SOPs & Policies.htm#4)

If you have questions regarding this information please contact Helene Lake-Bullock in the Office of Research Integrity (ORI) at Ph: 257-9428 or [hbullo@uky.edu](mailto:hbullo@uky.edu)