CC committee use:

**CC ID:**   **Recvd:**
**To SC:** **PI(s) notified:**
**Apprvd:** **Data delivered:**

|  |  |
| --- | --- |
| **Date:** |  |
| **Title of Proposed Project:**  |  |
| **Name of Principal Investigator (PI):**  |  |
| **PI Institution:** |  |
| **PI Email:** |  |
| **PI Telephone:** |  |
| **Proposed Collaborators: (name, institution, email)**  |  |
| **WOS Sites Holding Requested Data:** [ ]  City of Hope. Contact: Shiuan Chen schen@coh.org or Susan Neuhausen sneuhausen@coh.org [ ]  Columbia University. Contact: Mary Beth Terry mt146@columbia.edu or Rachel Miller rlm14@cumc.columbia.edu [ ]  George Mason University/University of Florida. Contact: Kevin Wright kwrigh16@gmu.edu or Carla Fisher carlafisher@ufl.edu [ ]  Georgetown University Medical Center/Uniformed Services University of the Health Sciences. Contact: Mary Beth Martin martinmb@georgetown.edu or Celia Byrne celia.byrne@usuhs.edu [ ]  Michigan State University/University of Cincinnati. Contact: Richard Schwartz schwart9@msu.edu [ ]  UCLA/University of Chile/Fox Chase Cancer Center/Silent Spring. Contact: Karin Michels k.michels@ucla.edu [ ]  University of Delaware. Contact: Kami Silk kamisilk@udel.edu [ ]  University of Massachusetts-Amherst/Baystate Medical Center. Contact: jjerry@vasci.umass.edu or Sallie Schneider sallie.schneider@baystatehealth.org [ ]  University of South Carolina. Contact: Daniela Friedman dfriedma@mailbox.sc.edu   |
| **Background/Significance.** 1-3 paragraphs providing the rationale for the project. Include citations. |
|   |
| **Specific Aims, Hypotheses, or Research questions.** |
|  |
| **Statistical Analysis Plan.** Describe the pertinent variables (independent and dependent variables with covariates), proposed modeling approach, and draft table shells. |
|  |
| **Are you requesting human or animal data?** If you are requesting both human and animal data, complete a separate form for the two requests. |
| [ ]  Animal [ ]  Human → **Do you already have human subjects approval to receive the data?** If you anticipate that this project will be determined to be *not human subjects research* or exempt from human subjects review, attach documentation to support this determination. Data will not be shared until evidence of IRB approval or waiver/exemption is provided.[ ]  No. Have not yet applied for human subjects approval. [ ]  Yes. Please attach the approval letter with this request. |
| **Anticipated Timeline.** |
|  |
| **Comments/Other.** |
|  |