

Request for Exclusion of Partial Study Data from EMR

Study data should only be excluded from a participant's EMR when the data is unusually sensitive, may be de-identified, or system is unable to process. Please complete the following form to request exclusion of specific study data. Form processing may take up to 8 business days, please plan accordingly.*

Provide the following information

Date submitted: **Projected start date:** **Projected end date**:**

IRB protocol number: **Fund/GL number:**

Personnel submitting request (first and last):

Principal Investigator (first and last):

PI email address:

Primary financial contact (first and last), if different:

Primary financial contact email address:

Primary financial contact phone number:

Entity: BWH ___ DFCI ___ MGH ___ Faulkner ___ Spaulding ___ McLean ___

Sponsor type: DHHS ___ Industry ___ Internal (sundry, other) ___ Non-profit ___ Other Federal Private Foundation ___ Public Foundation ___ State/local ___

Type(s) of service(s) requested

Clinical/Laboratory:

CPT Code(s):

Estimated project volume:

1. Does the study require tissue sample processing through BWH's PowerPath?
a. Yes ___ b. No ___

2. Does the study have de-identified samples that cannot be processed in a research CORE lab, and instead will be processed through the applicable institution's clinical lab?
a. Yes ___ b. No ___

3. Additional comments:

*Please, note: requests for "sensitive" approval, for all study data to be processed by masking the title of the study within the EMR and keeping the protocol #, must be approved within the study's IRB application. The Research Billing Review process will remain the same for "sensitive" studies. Click [here](#) more information.

** CLR # will close by the end of the quarter where the projected end date falls, unless otherwise instructed.

Research Compliance sign-off: _____ Date: _____

Research Controller sign-off: _____ Date: _____