Study Development and Study Support

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From protocol inception to IRB consultation and compliance management, we offer comprehensive support to shape robust study frameworks. Our expertise spans workflow design for recruitment and the creation of compelling communication materials to engage participants effectively.

Our suite of services encompasses study coordination, whether on-site or online, alongside meticulous recruitment coordination. We provide on-site assistance, routine metrics and reporting, and optimization of recruitment procedures, supplemented by tailored staff training to ensure seamless study execution.

ECRI is proud to be the managing team over patient related and regulatory operations for the following long-lasting projects and initiatives:

- Universal Consent for Biological Samples
- Athena Breast Health Network
- Wisdom Study
- Depression Grand Challenge
- Institute for Precision Health/ATLAS
- Center for Smart Health

If you are interested to collaborate with these programs, email us at ECRI@mednet.ucla.edu to schedule a discussion around the available opportunities.

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Drupal.jQueryUiFilter.globalOptions('accordion');