

### **OnCore**

OnCore is a clinical trials management system (CTMS) used at UCLA for the management of clinical trials in clinical research. The system maintains and manages planning, performing and reporting functions at the study and subject level along with participant contact information, tracking deadlines and milestones.

The use of OnCore enables the study team to set up and manage all protocols and subjects in one place, supports clinical research billing, simplifies data management and monitoring, improves patient recruitment and tracking and automates the flow of information between UCLA based systems including WebIRB, PATS and CareConnect.

Please see OnCore notifications for a better understanding of OnCore study staff roles and associated notifications.

OnCore is managed and maintained by the UCLA Clinical Research Informatics Systems (CRIS) team, which works closely with study teams to incorporate OnCore into their workflows for more efficient clinical research operations and centralized data management. The CRIS team works collaboratively with the <a href="Clinical Research Finance">Clinical Research Finance</a> team, Clinical Research Business Partners (CRBP) and Research Connect team members to provide a variety of data solutions and operation management tools for clinical research teams.

# **OnCore Training?**

- For system security and compliance, training is required <u>for access to OnCore</u>. Registration for online and inperson trainings is available through the <u>CareConnect Training Management System</u>
- To request an account and view training requirements for the different levels of access, view the research recommended training section available from the <u>UCLA CareConnect website</u>
- For training questions, contact <a href="mailto:crishelpdesk@mednet.ucla.edu">crishelpdesk@mednet.ucla.edu</a> or call 310-267-2273

#### OnCore Roles

WebIRB Role	BruinIRB Role	OnCore Role
Principal Investigator	Principal Investigator	UCLA Principal Investigator*
Fund Manager	Fund Manager	Fund Manager
Regulatory Coordinator	Regulatory Coordinator	Regulatory Coordinator
Study Contact	Primary Contact	Study Contact*
N/A	N/A	Billing Analyst
Study Coordinator	Research Coordinator	Study Coordinator (CRA)
Nurse Coordinator	Research Nurse	Nurse Coordinator
Research Assistant	Research Assistant	Research Assistant
Study Contact	Primary Contact	Recruiting Contact



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Data Manager	Data Manager	Data Manager
Co-Investigator	Co-Investigator	Sub-Investigator
Co-Principal Investigator	Co-Principal Investigator	Co-Principal Investigator
Statistician/Data Analyst	Statistician	Statistician
* These are the essential OnC	ore staff roles that must be included on a study of	coming from the IRB.

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