Bootcamp for New Investigators in Clinical Research

A series of four workshops on how to operationalize biologic clinical trials at GW and CNH.
An emphasis on available programs, templates, and resources.

Session 1. Thurs NOV 17, 12-2 pm (virtual)
Principal Investigator Responsibilities • Responsibilities for industry-sponsored or investigator-initiated studies, single/ multi-site: **Henry Kaminski** • Investigators rely on trained clinical research staff: **Aileen Chang** • Investigators need to recognize conflicts: **Laura Sigman**
Protocol & Study Frameworks • Cohort discovery & databases: **Hiroki Morizono** • Protocol Builder/ OnCore: **Richard Lush** • Investigational drug pharmacy-do-able: **Dorinne Mettle-Amuah**
Breakouts: Institutional solutions to challenges "How To Get Started as an Investigator in an Externally Sponsored Study" • **Radwa Aly**, GW OCR, Richard Lush GWCC Clinical Trials Office • **Jurran Wilson**, Patrick O'Keefe CNH CTR

Session 2. Thurs DEC 15, 12-2 pm (virtual)
Optimizing Roles on the Study Team • How the strengths of CRA and the investigator tie into the audit: **Radwa Aly** • Who is part of the Safety team? **Caitlin Joffe** • Optimizing stakeholder input: **Randi Streisand**
The clinical trial contract • Budgeting and Contracts: **Melanie Bossi** • Standard of care testing vs research costs: **Stephanie Bair** • Service biostatistics: **Qing Zeng**
Breakouts: Institutional Solutions to Challenges: *Who pays for what? Standard of care and protocols for escalation* • **Kiara Kalbart**, **Sheila Garrity** • **Bobbe Thomas**, **Stephanie Bair/ Khan**

For Zoom links and calendar invites, please contact Charlotte Hovland at cphovland@gwu.edu or gwsmhsresearch@gwu.edu

CTSI Clinical and Translational Science Institute at Children's National
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KL2-Special Interest Group Event