

Clinical Research Workshop: A Deep Dive into Informed Consent

March 23, 2023

8:30am - 2:00pm

Howard University - Louis Stokes Library

AGENDA

8:30 AM

Breakfast

9:00 AM

Introductions

Pre-T1

Informed Consent Process

- Panel - Coordinators (novice and experienced) - What are their pain points?
 - NOVICE PANEL
 - Niah Woods - Howard University
 - Bibiana Ateh - DC VAMC/ICR
 - Samantha Sterba - MHRI
 - EXPERIENCED PANEL
 - Jillian Turner - Howard University
 - Mikaley Bolden - Georgetown University
 - Stacy Malloy - MHRI

Post-Consent Studies - Jane Otado, Priscilla Adler

Florida Informed Consent Case Study Discussion - Petros Okubagzi

Remote/E-Consenting - Petros Okubagzi

Plain Language, Teach-back, and Guidance for Re-consent -
Shaunagh Browning

Role Play: Mini ICF Discussion - Claudia Gunawan

11:30 AM

Lunch

12:00 PM

Informed Consent Source Documentation -
Neha Mookuparambil, Senior QA auditor from Georgetown
University

12:30 PM

Special Populations/Vulnerable Populations - Florencia Gonzalez

- Medical Resource Document
 - Link:
(https://docs.google.com/document/d/1tfgaUSmMNw7DoGkJCyn1Stm152_K1XJF/edit?usp=sharing&oid=111336445613483281491&rtpof=true&sd=true)
- Guidance Document for Underinsured and Uninsured Study Participants

1:00 PM

Jeopardy Game

1:30 PM

Post - T2

Registration Link:

<https://redcap.link/GHUCCTSClinicalResearchWorkshop>

References:

- 1) Clinical Research Glossary - Health Literacy in Clinical Research (mrctcenter.org)
Using the Teach-back Toolkit – Teach Back Training
- 2) Culturally competent strategies for recruitment and retention of African American populations into clinical trials. J. Otado et al. (2015)
- 3) Satisfaction and perceptions of research participants in clinical and translational studies: An urban multi-institution with CTSA. P. Adler et al. (2020)

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