Office for the Protection of Human Subjects (OPHS) at Children's National - *Past Educational Sessions*

Session Title	Presenter(s)
Introduction to Research Regulatory Affairs	Maryann Rossi, PhD
Introduction to Human Subject Protections	Alicia Cook, BS, CIP; Maryann Rossi, PhD
Introduction to Study Design	Jim Chamberlain, MD
How to Read a Research Protocol	Caitlin Joffe, MBA, CCRP
Navigating the EMR to Collect Research Data	Juliana Lintner, BS; Sam Indresano, BS
Pathology and Lab Medicine Services	William Suslovic, BS
Clinical Research Unit	Elena Gibson, RN, CCRP
Overview of HHS and FDA Regulations	Alicia Cook, BS, CIP; Alavy Sos, MS
Reporting to the CNH IRB: What, When, and How	Kristen Breslin, MD, MPH
HIPAA Privacy Rule for Research	Megan Allen, MBA, CHPC, CHPS; Brittany Duah, MSP
Experts' Guide to Successful IRB Submissions - Part 1: Using IRBear 2.0	Alicia Cook, BS, CIP
Experts' Guide to Successful IRB Submissions - Part 2: Top 10 Mistakes to Avoid	Kathy Seabolt, CIP
Scientific Support Services for Human Research	Jermaine Lawson, MHA; Harley Little
Single IRB: Multi-site and Collaborative Research	Almarie Coleman, BS, CIP
Information, Comprehension and Voluntariness in Informed Consent/Parental Permission and Assent	Alicia Cook, BS, CIP; Maryann Rossi, PhD; Pedro Diaz
Informed Consent Process (Roleplay)	Alix Fetch, PhD; Emily Leibold, BSE, CCRP; Janiya Brooks
Ethical Considerations in Human Research and the Role of the Research Subject Advocate	Vanessa Madrigal, MD, MSCE, HEC-C
Research Documents	Caitlin Joffe, MBA, CCRP
Research Monitoring and Quality Assurance	Caitlin Joffe, MBA, CCRP
Working with Your Biostatistician	James Bost, MS, PhD
Professional Development for Clinical Research Coordinators	Sarah Anderson; Alix Fetch, PhD; Emily Leibold, BSE, CCRP; Michael Bernardo, MS
When Does a Quality Improvement Project Become Human Subjects Research?	Kristen Breslin, MD, MPH; Kathy Seabolt, CIP
Requesting Emergency Use of an Investigational New Drug (IND)	Marissa Horrigan, Pharm D; Kristen Breslin, MD, MPH
Is Informed Consent Always Needed for Research?: Waivers and Alterations Explained	Lindsay Ropchock, JD, CIP
IRB Clinical Trial Review and Inspection: Readiness and Conduct	Caitlin Joffe, MBA, CCRP