As society moves away from a paternalistic view of protecting research participants, many are revisiting the ethics and utility of payment for research participation. The goal of this workshop is to explore the role of paying people for their participation, the impact of payment, as well as the challenges and considerations for how payments should be made. Session 1 will focus on ethical and theoretical considerations. Session 2 will explore the practical challenges and implementations.

### AGENDA

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<tr>
<td>10:00 AM – 10:05 AM</td>
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| 10:05 AM – 1:00 PM | **Session I: Ethical and Theoretical Considerations**  
**Moderator:** Neal Dickert, M.D., Ph.D; Associate Professor, Emory University School of Medicine, Division of Cardiology, Emory University Rollins School of Public Health, Department of Epidemiology, Emory Health Services Research Center, Emory Center for Ethics  
**Advancing Justice, Equity, and Inclusion through Payment for Research Participation**  
Luke Gelinas, Ph.D; Sr. Chair Director, Advarra  
**What Do IRBs and Investigators Think About Payment for Research Participation?**  
Emily A. Largent, J.D., Ph.D, R.N.; Emanuel and Robert Hart Assistant Professor of Medical Ethics, Department of Medical Ethics and Health Policy, University of Pennsylvania Perelman School of Medicine  
**Participants’ Perspectives I:**  
Amie Devlin, Ph.D, M.P.H., M.A.; Epidemiologist, Stratis Group – BluePrint Orphan  
**Participants’ Perspectives II:**  
Alexandra Collins, Ph.D; Assistant Professor, Brown University School of Public Health  
**Participants’ Perspectives III:**  
Dustin C. Krutsinger, M.D., M.S.; Assistant Professor of Medicine, University of Nebraska Medical Center in Omaha |
| 11:30 AM – 11:45 AM | Break                                                                                                                                 |
|                  | **Who Participates in Phase I Clinical Trials and What Are Their Motivations for Participation?**  
Jill A. Fisher, Ph.D; Professor of Social Medicine, Center for Bioethics, University of North Carolina at Chapel Hill  
**Are Phase I Clinical Trial Participants Workers and if so, What Rights Should They Have?**  
Roberto Abadie, Ph.D; Assistant Professor Department of Anthropology, University of Nebraska-Lincoln |

## AGENDA

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| 1:00 PM – 1:50 PM | Panel Discussion
Includes Special Guest: Jill Feldman; Lung Cancer Patient and Advocate, Co-founder EGFR Resisters |
| 1:50 PM – 4:00 PM | Lunch                                                                    |
| 1:50 PM – 4:00 PM | Session II: Practical Challenges and Implementations
Moderator: Quincy Byrdsong, Ed.D, CIP, CCRP; Vice Provost for Health Affairs, Lipscomb University |
|                   | Are Payments All the Same and Should We Always Offer the Same Payment to Participants in the Same Study?
David Borasky, M.P.H.; Vice President, IRB Compliance, WCG IRB |
|                   | Addressing the Financial Burden of Clinical Research Participation
Ivy R. Tillman, M.S., CCRC, CIP; Director of Research Operations, Mayo Clinic |
|                   | The Logistics of How We Pay People for Research Participation – Equity Considerations
Christine Ritchie, M.D., M.S.P.H.; Professor of Medicine, Harvard Medical School and Director, Mass General Hospital Mongan Institute Center for Aging and Serious Illness |
|                   | Paying For Participating in Social-Behavioral Studies – Practical and Ethical Considerations
Celia B. Fisher, Ph.D; Marie Ward Doty University Chair in Ethics, Professor of Psychology, Director, Center for Ethics Education and the HIV/Drug Abuse Prevention Research Ethics Institute, Fordham University |
|                   | It’s Not All about the Benjamins– Other Hurdles Presented by Research Participation, and Why Payment Won’t Resolve Them
C.K. Wang, M.D.; Chief Medical Officer, COTA, Inc. |
|                   | Panel Discussion
Includes Special Guest: Jill Feldman; Lung Cancer Patient and Advocate, Co-founder EGFR Resisters |