

PROGRAM

11:00 – 11:05 Introduction to MISCH Hub *Prof. Julie Simpson (MISCH Director)*

11:05 – 11:10 Goals of the Clinical Trial Coordinators Network Assoc Prof. Adam Deane (MISCH Clinical Trials Node Leader) Katie Ozdowska (MISCH Clinical Trials Officer)

11:10 – 11:15 Clinical Trial Network History & Success Sofia Sidiropoulos (Dept. Critical Care – Clinical Trial Manager)

11:15 – 11:25 Investigator Initiated Trial Toolkit Michelle Iddles (Melbourne Academic Centre for Health – Manager)

11:25 – 11:35 UoM Clinical Trial Governance Victoria McMorran (Office of Research & Ethics Integrity – Clinical Trials Governance Officer)

11:35 – 11:45 Legal Services Contracts & Agreements Johannah Planache (MDHS – Clinical Trials Contracts Officer)

11:45 – 11:50 HeSANDA Katie Ozdowska (MISCH Clinical Trials Officer)

11:50 – 12:00 Open Floor





Twitter @MISCHHub



Goals of the Clinical Trial Coordinators Network

- Resource and connect clinical trial coordinators across the University
- Communicate changes to University policy, guidelines, and highlight new developments
- MISCH Hub to host a repository of clinical trial resources
- Promote professional development
- Support new staff and retain experienced staff & advertise job vaccancies



The Health Studies Australian National Data Asset HeSANDA

- Led by the Australian Research Data Commons (ARDC)
- 9 Nodes around Australia encompassing 72 research organisations
- UoM is part of the MACH Clinical Trial Consortium Node partnership
- Developing national infrastructure to support the sharing and reuse of health research data.



Project aims & outcomes

- Stakeholder engagement working with MACH HREC's and institutions to deliver awareness around HeSANDA
- Building upon existing culture and policies required to ensure it's beneficial for the research and wider community
- Ensure trials follow standard trial metadata definitions across
 MACH partners
- Improving environments for metadata storage and transfer
- Advocacy for future trials to contribute to the HeSANDA database and to incorporate data sharing intentions from the start of trial development



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