New/Transfer PI Orientations: Handouts and guidance

1. Checklist (v.4)
2. PI Responsibilities (CCI Policy)
3. Registration of Clinical Trial w/International Committee of Medical Journal Editors
4. CCI/IRB: Guidelines & Policies – Table of Contents
5. Guidance: Electronic Storage of Study Data and Documents
6. Clinical Trial Study Document Checklist
7. CRP Guidelines for Document Retention and Destruction in Clinical Research
8. CCI Policy for Modifications: Exceptions and Deviations
9. Flowchart for Unanticipated Problems Involving Risk Reporting
10. EQuIP Guidance: Access to Online Medical Records for non-CHB Personnel
11. CRP: What can the CRP do for you?
12. GCRC (CTSU) Services and Contact Handout
13. Translational Research Program (TRP): Regulatory Resources
14. EQuIP Services Handout

15. Offer - Regulatory Binder Template
16. IF NEEDED, offer/lend copy of CRC Guide's to Coordinating Clinical Research

If possible, go through internet links on laptop, including:

1. Research Link from internal website (4 research squares)
2. Link to Informed Consent Library
3. CCI/IRB:
    a. Training Certificates
    b. IRB Member Reviewer Worksheets
    c. Policies
    d. Protocol Database Access
4. CRP: Guidelines and Services
5. EQuIP: Templates and Guidance