

Suggested Training for Clinical Researchers at UCSF

| # | Topic | Resource | Notes | Behavioral | Observational | Interventional | Cert/Lead/Documentation Provided | Approx. length of time to complete (min) |
|---|---|---|---|------------|---------------|----------------|----------------------------------|--|
| | | | | | | | | |
| New Hire and Onboarding Resources (Recommended Week 1-2) | | | | | | | | |
| 1 | HIPAA 101 | https://training.ucsf.edu/ | HIPAA 101 - Privacy and Security for New UCSF Faculty, Staff, Trainees, Students and Volunteers | R | R | R | Y | 20 |
| 2 | HIPAA and Data Security for Researchers (IRB) | https://irb.ucsf.edu/hipaa | https://irb.ucsf.edu/electronic-data-security | R | R | R | | 30 |
| 3 | CITI Human Subjects Training | http://irb.ucsf.edu/citi-human-subjects-training | Required training for all human research studies | R | R | R | Y | 240 |
| 4 | CITI GCP | https://www.citiprogram.org/ | Good Clinical Practice training is usually optional but highly recommended | R | R | R | Y | 360 |
| 5 | Add CRC to study in systems | Add to IRB applications (in IRIS), OnCore (clinicaltrials@ucsf.edu) and APEX study builds | https://ucsf.service-now.com/ess/ | R | R | R | | 45 |
| 6 | Add CRC to study documents | Add to delegation log in regulatory binder and document training | Be sure all studies track staff responsibilities and training | R | R | R | | 30 |
| 7 | Add to CRC ListServ | https://irb.ucsf.edu/clinical-research-coordinators-council#listserv | Also contact department/division admin for other distribution lists | R | R | R | | 10 |
| 8 | Occupational Health Screening | https://occupationalhealthprogram.ucsf.edu/home | You or your supervisor must make an appointment to have health screening clearance prior to exposure | R | R | R | Y | 10 |
| 9 | Review IRB Website | http://irb.ucsf.edu/ | Institutional Review Board (IRB); previously known as 'CHR'; selected links below | R | R | R | | 30 |
| 10 | IRIS Training (IRB) | https://iris-help.ucsf.edu/irb-iris | IRB training videos for online application system | C | C | C | | 60 |
| 11 | Infection Control Training (UCLearning) | https://training.ucsf.edu/ | Infection Control Training for Ancillary Staff | C | C | R | Y | 30 |
| 12 | Annual Safety Training (UCLearning) | https://training.ucsf.edu/ | Annual Safety Training for Staff | C | C | R | Y | 60 |
| 13 | Bloodborne Pathogen Training (UCLearning) | https://training.ucsf.edu/ | Bloodborne Pathogens (Campus) | C | C | R | Y | 45 |
| 14 | Safe Shipping (UCLearning) | https://training.ucsf.edu/ | Safe Shipping of Biohazards/Dry Ice | a | a | a | Y | 30 |
| 15 | Review The HUB Website and Training List | http://hub.ucsf.edu/ | Clinical Research Resource HUB (lots of info for CRCs) | R | R | R | | 15 |
| 16 | OnCore | https://trialactivation.ucsf.edu/oncore-account-creation-training | Complete Intro course on UC Learning Ctr to request access or email OnCore@ucsf.edu. | a | a | a | Y | 120 |
| 17 | APeX CRC Knowledge Bank and Training | http://myapex.ucsf.edu/researchcrc | UC Learning Center > APeX > 'Non-Clinical Staff' > 'Clinical Research Coordinator' > 'CRC' | a | a | a | R | 120 |
| Core Training (Recommended Weeks 2-6) | | | | | | | | |
| 18 | Role of the CRC (IRB) | http://irb.ucsf.edu/responsibilities-pis-and-crcs | Summary of the responsibilities of the PI and CRC | R | R | R | | 10 |
| 19 | Orientation (CTSI) | https://crs.ucsf.edu/crc-training | Sign up for the CRC Orientation Course | C | C | R | | 60 |
| 20 | Informed Consent Info (IRB) | http://irb.ucsf.edu/obtaining-and-documenting-informed-consent | IRB guidance on the informed consent process | R | R | R | | 30 |
| 21 | Informed Consent Training (CTSI) | https://crs.ucsf.edu/crc-training | Sign up for CRC 101 Training Course | C | C | C | | 90 |
| 22 | Research Integrity Training | http://ori.hhs.gov/TheResearchClinic | Entertaining interactive video for training research staff | C | C | C | | 60 |
| 23 | Introduction to The Regulatory Binder (HUB) | https://hub.ucsf.edu/virtual-regulatory-binder | Guidance on creation and maintaining essential documents | a | C | R | | 30 |
| 24 | Research Tools and Enrollment Logs (IRB) | http://irb.ucsf.edu/research-tools-and-checklists | Templates for study logs and other study tools | R | R | R | | 30 |
| 25 | Amendments and Version Control (IRB) | http://irb.ucsf.edu/modification | Guidance regarding protocol amendments and modification applications | C | C | C | | 10 |
| 26 | Study Start-up and Billing (CTSI/OCTA) | https://crs.ucsf.edu/crc-training | Sign up for CRC 103 Training Course | C | C | C | | 90 |
| 27 | Study Start-up Checklist (HUB) | https://hub.ucsf.edu/clinical-research/setup-study | Steps to begin new clinical from receipt of protocol to enrollment of 1st subject | C | C | C | | 30 |
| 28 | IRB and Safety Reporting (CTSI) | https://crs.ucsf.edu/crc-training | Sign up for CRC 102 Training Course | C | R | R | | 90 |
| 29 | Budgets and Coverage Analysis (HUB) | http://hub.ucsf.edu/ca-budget-billing | Describes the importance of compliant clinical research billing practices | C | C | R | | 30 |
| 30 | Post-approval Reporting Summary (IRB) | https://irb.ucsf.edu/sites/hrpp.ucsf.edu/files/post-approval-reporting-summary-sheet-%20%286-1-20%29.pdf | A summary sheet of all AE/SAE, safety, new IB, DSMB, protocol incident and violation reporting | R | R | R | | 30 |
| Supplemental or Advanced Training Resources for CRCs | | | | | | | | |
| 31 | Data Management for CRCs (CTSI) | https://crs.ucsf.edu/crc-training | Sign up for CRC 106 Training Course | R | R | R | | 30 |
| 32 | Audit Readiness | https://crs.ucsf.edu/crc-training | Sign up for CRC 201 Training Course | a | C | R | | |
| 33 | Participant incentive policies (IRB) | http://irb.ucsf.edu/research-subject-payments | Guidance regarding subject payments | a | a | a | | 10 |
| 34 | Policies for Petty Cash Handlers (UCLearning) | https://training.ucsf.edu/ | Useful if you handle gift cards for research participants | a | a | a | Y | 60 |
| 35 | Subject Injury Program | http://irb.ucsf.edu/treatment-and-compensation-injury | Policy on reporting subject injury (AE as a result of study participation) | a | C | R | | 10 |
| 36 | Recruitment (IRB and HUB) | http://irb.ucsf.edu/recruitment | https://recruit.ucsf.edu/ | C | C | C | | 15 |
| 37 | IND Process (CTSI) | https://crs.ucsf.edu/crc-training | https://hub.ucsf.edu/ind-development-process | C | a | C | | 10 |
| 38 | IDS Pharmacy Information | http://ids.ucsf.edu/ | Log-in using your MyAccess account | a | a | C | | 10 |
| 39 | Protocol Development (HUB) | https://hub.ucsf.edu/protocol-development | Basic information about protocol organization and development | a | a | a | | 10 |
| 40 | CRS Procedures and Budget Estimate | https://accelerate.ucsf.edu/research/crs | For studies that use CRS, PCRC or other CTSI services (look up by location) | a | a | a | | 10 |

For any questions about this list, please email:

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CRC Training and Resource Advisor, CTSI

KEY:
R = required;
C = recommended;
a = as needed, or if applicable