**Study Startup** *100 Series*

UCSF 101 Study Startup Process

UCSF 102 Coverage Analysis

UCSF 103 Faculty Oversight Committee for Research (FOCR)

UCSF 104 Patient Care Manager (PCM) Communications

DOM 105 Safety Committee Submissions

DOM 106 New Products Committee Submission

DOM 107 CMS Submissions

DOM 108 APeX Study Build Request (Maintenance requests)

DOM 109 OnCore Clinical Trial Management Systems

**Regulatory** *200 Series*

UCSF 201 Regulatory Binder

DOM 202 Source Documents

DOM 203 FDA 1572

DOM 204 Financial Disclosure

DOM 205 Delegation Log Completion

DOM 206 Documentation of Training

**Clinical Research Coordinator Orientation** *300 Series*

DOM 301 CRC Orientation

DOM 302 Informed Consent

DOM 303 Recruitment Methods

DOM 304 Atomic Web

DOM 305 Scheduling Research Visits

**Study Management** *400 Series*

DOM 401 Study Manual of Procedures (MOP)

DOM 402 Participant Accrual (APeX, OnCore)

DOM 403 Participant Reimbursement

DOM 404 Billing Review in APeX

DOM 405 Biological Specimen Shipping

DOM 406 Investigational Product Accountability

DOM 407 Investigational Transfer

DOM 408 Protocol Amendments (ICF Revisions, CA/budget amend/ training)

DOM 409 Record Retention and Destruction

DOM 410 Study Closure

**Compliance** *500 Series*

DOM 501 CHR Initial Submission

DOM 502 Continuing Review (Amendments)

DOM 503 Safety Reporting

DOM 504 Note-to-File

DOM 505 Violations

DOM 506 Data Security (HIPAA)

DOM 507 Monitoring

DOM 508 Audits

DOM 509 Equipment Calibration

**General Administrative** *600 Series*

DOM 601 Approval and Review of SOPs

DOM 602 Sub-Site Management

DOM 603 Effort Tracking