Templates & Resources

Note: Some links may require logging in with your AD credentials.

- <u>Clinical Study Management Process</u>
- <u>Creating Study Design</u>
- Glossary of Lay Terms for Use in Preparing Protocols for the IRB
- ICH Essential Documents for the Conduct of a Clinical Trial
- IRB Assurance Letter/FWA Letter
- Personal Vehicle Usage for UCSD Business (see Blink page for Insurance Requirements and Liability Issues)
- Protocol Overview Fact Sheet Template
- Quick Reference Guide Clinical Trial and Investigator-Initiated Agreements
- Radiation Risk Calculator
- Registration of Clinical Trials in a Public Registry for Publication
- <u>Research Acronyms & Definitions</u>
- <u>Storage Facilities</u>
- Velos: UCSD's Clinical Trial Management System
- Sample Standard Operating Procedure
- Standard Operating Procedure Template
- <u>Regulatory Binder Template</u>

Last updated: 17 Sep 2019

Source URL: https://compass.ucsd.edu/templates-resources

Drupal.jQueryUiFilter.globalOptions('accordion');

<u>Clinical Research Billing | CRB</u>

Human Research Protections Program | HRPP

Office of Coverage Analysis Administration | OCAA

Office of Contract & Grant Administration | OCGA