Welcome

- **Research in Response to COVID-19**

  In an effort to eliminate apparent immediate hazards to subjects (45 CFR 46.108(3)(iii)) and the Stanford community, all in-person visits for non-essential human research activities must be postponed, while preserving critical activity on interventional treatment trials or critical clinical research. In light of this urgency around COVID-19, *Stanford’s IRB is not requiring prior review of some protocol changes*, e.g., implementing video/phone visits, mailing of study medications, or other changes designed to protect participant safety. Understanding that resources must be prioritized, please track these changes and file a modification in [eProtocol](https://researchcompliance.stanford.edu) when your schedule allows. The IRB is available to assist you with these modifications.

  Please refer to [Stanford Health Alerts](https://researchcompliance.stanford.edu) and [Stanford Health Care](https://researchcompliance.stanford.edu) webpages. Note that the process for [Emergency Use](https://researchcompliance.stanford.edu) of an experimental drug or device remains the same.

  Contact IRB Education via [email](https://researchcompliance.stanford.edu) / (650) 724-7141 or your [Panel Manager](https://researchcompliance.stanford.edu) for questions.

- **Survey Results (2019-2020 Panel Year Q1 Results)**

- **What’s New**
  - **New** RCO is moving to 1705 El Camino Real, Palo Alto CA 94306 / SU ID Mail 5579 (effective 1/1/2020)
  - **New** Common Rule effective January 21, 2019
  - Informed consent templates have been updated to include the new requirement to provide key information as a concise summary at the beginning of the consent form, the new basic element of consent regarding future use, and the new additional elements of consent as applicable. (The concise summary is not required for Minimal Risk or Non-medical consents.)
  - **New** Exempt Form in eProtocol (video)
  - eProtocol is rolling out a new user interface (UI) on the updated Exempt form that includes the new Common Rule Exempt categories. While the new UI has a different appearance, the application questions have not changed. The Exempt form is the only IRB form using the new UI at this time.
  - **New** The revised Common Rule requires that for any clinical trial conducted or supported by a Common Rule department or agency, one consent form must be posted on a publicly available federal website after recruitment closes and no later than 60 days after the last study visit. The consent form must have been used to enroll subjects in order to satisfy this new provision. Click [here](https://researchcompliance.stanford.edu) for details.
  - [Sample eProtocol Template Forms](https://researchcompliance.stanford.edu)
  - Two new eProtocol forms, Single Patient IND and Humanitarian Use Device (HUD)
  - European Union General Data Protection Regulation (GDPR) is now in effect. [Learn more](https://researchcompliance.stanford.edu)
  - [Single IRB (sIRB)](https://researchcompliance.stanford.edu)
    - Relying on a sIRB and sIRB SOP
    - NCI CIRB - the NCI Central IRB Initiative
  - ClinicalTrials
Revised Intake Process for Industry Clinical Trials

Changing policies impact NIH-funded studies involving human subjects

In your proposal application packet, be sure to complete the ?R&R Other Project Information? form page before you complete the ?PHS Human Subjects and Clinical Trials Information? form page as the first will populate the second. Please view this 9 min You Tube video: PHS Human Subjects and Clinical Trial Information Form Walk-through.

ClinicalTrials.gov

CRQ’s Clinical Trials.Gov resources page

Quick links for researchers

Getting started

- For Researchers
- Consent Process
- Human Subjects Research (HSR) Determination application is available in eProtocol. Attach this completed form. (More Information)

Consent Templates, Forms, eProtocol attachments

- Medical - consents, minimal risk samples, short form
- Nonmedical - consents, parental permission
- Medical & Nonmedical - assent, phone scripts and screens, etc.
- eProtocol Required Attachments (VA, international research, etc.)
- Sample eProtocol Applications

FAQs

Basics | Training | Protocol Management | Single IRB | Genomic Data Sharing | Cadavers or Deceased Individuals | Certificate of Confidentiality | GDPR | Other/Special Circumstances

- About the IRB (the Administrative Panels on Human Subjects Research)
- Charge to the IRBs

Source URL: https://researchcompliance.stanford.edu/panels/hs

Links
[1] https://eprotocol.stanford.edu/
[5] mailto:irbeducation@stanford.edu
[10] https://eprotocol.stanford.edu/irb