Welcome

Research in Response to COVID-19


- **COVID IRB Resources** [2]

**COVID-19 Clinical Research Review Panel**

Researchers considering any COVID-19 clinical research, including studies that require Institutional Review Board (IRB) approval as well as quality improvement projects, are required to complete a short survey at [COVID-19 Clinical Research Review](https://researchcompliance.stanford.edu) [3]. For more information, click [here](https://researchcompliance.stanford.edu) [2].

- Contact the Research Participation team at [EngageParticipants@stanford.edu](mailto:EngageParticipants@stanford.edu) [4] to list your study in the [Stanford COVID-19 Study Directory](https://researchcompliance.stanford.edu) [5].

Satisfaction Surveys

- Satisfaction Survey Results Panel Year 2020-2021 [6]
- Satisfaction Survey Results Q1 Panel Year 2021-2022 [7]

What's New

- New *Gene Transfer eP Supplemental Questionnaire* [8] (required for human gene transfer studies)
- Exempt Information Sheet template for Non-Medical/Social Behavioral studies [HERE](https://researchcompliance.stanford.edu) [9].
- Training videos posted [HERE](https://researchcompliance.stanford.edu) [10].
- New Common Rule effective January 21, 2019 [12].
- Sample eProtocol Template Forms [13].
- European Union General Data Protection Regulation (GDPR) is now in effect. [Learn more](https://researchcompliance.stanford.edu) [16].
- Single IRB (sIRB) [17]
- ClinicalTrials
  - Changing policies impact NIH-funded studies involving human subjects [18] 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](https://researchcompliance.stanford.edu) [19].
- ClinicalTrials.gov [20]
  - CRQ's [Clinical Trials.Gov resources page](https://researchcompliance.stanford.edu) [21]
Quick links for researchers

Getting started

- For Researchers [22]
- Consent Process [23]

Consent Templates, Forms, eProtocol attachments

- Medical - consents, minimal risk samples, short form [26]
- Nonmedical - consents, parental permission [9]
- Medical & Nonmedical - assent, phone scripts and screens, etc. [27]
- eProtocol Required Attachments [13] (VA, international research, etc.)
- Sample eProtocol Applications [28]

FAQs [29]


- About the IRB [39] (the Administrative Panels on Human Subjects Research)
- Charge to the IRBs [40]

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

Links
[1] https://cardinalrecovery.stanford.edu/research/research-recovery/
[3] https://app.smartsheet.com/b/form/324e5813005e465cad070e46fd5de753
[4] mailto:EngageParticipants@stanford.edu
[8] https://stanfordmedicine.box.com/shared/static/zlr02afphvcwgsr0cez3px13i9igf58x.docx
[14] https://eprotocol.stanford.edu/
[17] https://researchcompliance.stanford.edu/panels/hs/forms-for-researchers/singleirb
[18] https://stanfordmedicine.box.com/shared/static/umvuw8n8cms8o7db4t7gpot3x41tgbf.pdf
[19] https://www.youtube.com/watch?v=nz9NWFhYOG8&amp;list=PLOEUwSnjvqBJeHcb4yai7_fDnFZFPFEmQK&amp;index=1