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** [3]. For more information, click [here](https://researchcompliance.stanford.edu) [2].

- Contact the Research Participation team at EngageParticipants@stanford.edu [4] to list your study in the **Stanford COVID-19 Study Directory** [5].

Satisfaction Surveys

- **Satisfaction Survey Results Panel Year 2020-2021** [6]
- **Satisfaction Survey Results Q4 Panel Year 2020-2021** [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 [9].
- Training videos posted HERE [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 [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 [19].
- **ClinicalTrials.gov** [20]
  - CRQ's Clinical Trials.Gov resources page [21]
## Quick links for researchers

### Getting started

- **For Researchers** [22]
- **Consent Process** [23]
  - Human Subjects Research (HSR) Determination application is available in [eProtocol] [14]. Attach this completed form [24]. ([More Information] [25])

### 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]

- **Basics** [30] | **Training** [31] | **Protocol Management** [32] | **Single IRB** [33] | **Genomic Data Sharing** [34] | **Cadavers or Deceased Individuals** [35] | **Certificate of Confidentiality** [36] | **GDPR** [37] | **Other/Special Circumstances** [38]

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

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**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
[7] https://stanfordmedicine.box.com/shared/static/1i2v6kmhtcfuq8rq5855tmyx1jveqjak.pdf
[8] https://stanfordmedicine.box.com/shared/static/zlr0afphvcwgrs0cez3px13i9igf58x.docx
[14] https://e.protocol.stanford.edu/
[17] https://researchcompliance.stanford.edu/panels/hs/forms-for-researchers/singleirb
[18] https://stanfordmedicine.box.com/shared/static/umvvwu8n8cms8o7db4t7gpot3x41tgft.pdf
[19] https://www.youtube.com/watch?v=nz9NWFhYOG8&amp;list=PLOEUwSnjvqBJeHcb4yai7_fdN4ZFPEmQK&amp;index=1