Training

Education and training are provided to all individuals involved with the Human Research Protection Program (HRPP). The HRPP Policy Manual [1] specifies education requirements for IRB members, IRB staff and research personnel.

**eProtocol Training**

eProtocol training sessions (slides [2]), for new and less experienced users, are provided by the Research Compliance Office. Click [here to register] [3] for an upcoming eProtocol training session.

Stanford provides access to required training through the interactive online Collaborative Institutional Training Initiative (CITI [4]) Course: The Protection of Human Research Subjects. ([more information] [5])

**IRB Member Education**

View past training presentations, materials and handouts [6] from previous panel meetings.

**Other Resources**

- [Presentations] [7] provided to the research community and IRB
- [FAQs] [8]
- eProtocol FAQs [9]
- [Spectrum] [10] - Stanford/Packard Center for Translational Research in Medicine
- [CCTO] [11] - Stanford Cancer Clinical Trials Office

**Additional training:**

As well as the human subjects training for researchers required by the IRB, there might be training required by other groups, programs, or departments. Researchers should see Training [12] on the Dean of Research (DoR) [13] website, or refer to their department to identify needed education.

**Questions?**

Contact [IRB Education] [14] or call (650) 724-7141

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**Source URL:** https://researchcompliance.stanford.edu/panels/hs/forms/training

*Links*

[2] https://stanfordmedicine.box.com/shared/static/gixxa93hkkbg1s3n87jqvg0hfenmhj4d.pptx
[14] mailto:IRBEducation@stanford.edu?subject=IRB%20Education%20query