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**

Click [HERE][2] to register for an upcoming **eProtocol Training Session**.

eProtocol training sessions (slides [3]), for new and less experienced users, are provided by the Research Compliance Office.

Please review the following videos for information on:

- Who/What is the IRB? [4]
- Getting Started - What is Needed and eProtocol Basics [5]
- How to Submit a Continuing Review or Modification in eProtocol [7]

Stanford provides access to required training through the interactive online Collaborative Institutional Training Initiative (CITI [9]) Course: The Protection of Human Research Subjects. ([more information][10])

**IRB Member Education**

View past training presentations, materials and handouts [11] from previous panel meetings.

**Other Resources**

- [Presentations][12] provided to the research community and IRB
- [FAQs][13]
- [eProtocol FAQs][14]
- [Spectrum][15] - Stanford/Packard Center for Translational Research in Medicine
- [CCTO][16] - 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 [17] on the Dean of Research (DoR) [18] website, or refer to their department to identify needed education.

**Questions?**