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 sessions (slides [2]), for new and less experienced users, are provided by the Research Compliance Office:

<table>
<thead>
<tr>
<th>DATE</th>
<th>TIME</th>
<th>LOCATION</th>
<th>REGISTER</th>
</tr>
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<tbody>
<tr>
<td>October 22, 2019</td>
<td>2:30-4:00 PM</td>
<td>Turing Auditorium</td>
<td>Click here [3]</td>
</tr>
<tr>
<td>January 15, 2020</td>
<td>2:30-4:00 PM</td>
<td>Turing Auditorium</td>
<td>Click here [4]</td>
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CITI

Stanford provides access to required training through the interactive online Collaborative Institutional Training Initiative (CITI [5]) Course: The Protection of Human Research Subjects. (more information [6])

IRB Member Education

View past training presentations, materials and handouts [7] from previous panel meetings.

Other Resources

- Presentations [8] provided to the research community and IRB
- FAQs [9]
- eProtocol FAQs [10]
- CCTO [12] - 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 [13] on the Dean of Research (DoR) [14] website, or refer to their department to identify needed education.

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

Links
[2] https://stanfordmedicine.box.com/shared/static/gixxa93hkkg1s3n87jqvg0hfenmhj4d.pptx
[8] https://researchcompliance.stanford.edu/panels/hs/forms/training/presentations
[15] mailto:IRBEducation@stanford.edu?subject=IRB%20Education%20query