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>
</thead>
<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>
</tr>
</tbody>
</table>

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?