Highly Recommended Resources for IRB Members

[The Belmont Report](http://www.fda.gov/ohrms/dockets/ac/05/briefing/2005-4178b_09_02_Belmont%20Report.pdf)

[CITI, IRB Member Training Instructions](https://researchprotections.appstate.edu/sites/researchprotections.appstate.edu/files/CITI%20Training%20Instructions%202014.pdf)

* **[Researchers normally take Human Subjects training but there is a course specific to IRB members.  This link has directions on how to register for this course.  The link to CITI is in the Supplementary Resources section.](https://researchprotections.appstate.edu/sites/researchprotections.appstate.edu/files/CITI%20Training%20Instructions%202014.pdf)**

[45 CFR 46](https://www.hhs.gov/ohrp/sites/default/files/ohrp/policy/ohrpregulations.pdf)

* **[(Common Rule- the current regulations that we follow)](https://www.hhs.gov/ohrp/sites/default/files/ohrp/policy/ohrpregulations.pdf)**

[ASU’S IRB Policies & Guidelines](https://researchprotections.appstate.edu/human-subjects/irb-policies-guidelines)

[IRBIS](https://appstate.myresearchonline.org/irb/index_auth.cfm)

Supplementary Resources

[CITI Program Website](https://www.citiprogram.org/index.cfm?pageID=14&languagePreference=English&region=1)

[Free IRB Newsletter Quorum Review](http://www.quorumreview.com/)

[New ‘Common Rule’ for Research: Federal Register](https://www.federalregister.gov/documents/2017/01/19/2017-01058/federal-policy-for-the-protection-of-human-subjects) Released 1/18/2017

* **The Final Rule above goes into effect in 2018.**

[FDA’s webpage on the IRB](http://www.fda.gov/AboutFDA/CentersOffices/OfficeofMedicalProductsandTobacco/CDER/ucm164171.htm)