Here's a list of Learning Shots

The University of Virginia in Charlottesville, VA, has created brief, on-line, educational sessions for the human subjects research community, including investigators and clinical trial staff.
Some of the "Learning Shots," located at http://www.virginia.edu/vpr/irb/hsr/education_online.html include these topics:

- HIPAA & Waiver of Consent Part 1: Terminology (15 min);
- HIPAA & Waiver of Consent Part 2: Privacy Plan (15 min);
- Safety Reporting Part 1: Adverse Events (8 min);
- Safety Reporting Part 2: Protocol Violations and Unanticipated Problems (12 min);
- Use of consent and assent forms to enroll minors (12 min);
- Use of short forms for non-English speaking subjects (8 min);
- Recruitment, advertising and HIPAA (10 min);
- Process for modifying IRB-approved protocols (7 min);
- The informed consent process for clinical research (11 min);
- Tips for writing better research consent forms (13 min);
- Using bullets and tables in your research consent form (4 min);
- IRB-HSR protocol submission requirements for international research (13 min);
- Continuing review and five year updates (8 min);
- Post-approval monitoring (PAM) program overview (7 min);
- Source documentation (14 min);
- Clinical researcher roles and responsibilities (14 min);
- New protocol review process for non-scientists (11 min);
- Continuations (9 min);
- Modifications (7 min).