2017 Human Research Protection Program (HRPP)
Brown Bag Series
First Tuesday of each month at noon

Topics subject to change and/or reorder

**February 7**, Auditorium C: Summary and discussion of the revisions to the Common Rule

**March 7**, Auditorium B: External IRBs, Dartmouth as IRB of Record, Single IRB (NIH requirement)

**April 4**, Auditorium B: Data Safety Monitoring and Reporting requirements

**May 2**, Auditorium C: Welcome Leigh Burgess, VP Research Operations at D-H. Leigh will describe her role and provide an update on new requirements for clinical trial.gov

**June 6**, Auditorium B: Scientific Review Process

**July 11th (second Tuesday)**: Borwell 658W: Quality Improvement

**August 1**, Auditorium B: Global consent for research (CTSA consortium)

**September 5**, Auditorium C: Written consent form / teach back

**October 3**, Auditorium B: Recruitment, prioritize research studies

**November 7**, Auditorium B: Investigational drug management

**December 5**, Auditorium B: Significant Financial Interest / Conflict of Interest review