ADAPT
Addressing Disparities in Asian Populations through Translational Research (ADAPT)

Course description: The ADAPT seminars aim to raise awareness of Asian health disparities and facilitate community-engaged translational research that targets the health of Asian-Americans. These presentations are for the research and clinical communities, public health and policy makers, and community leaders.

1. Together Strengthening the Health of Chinatown: Advancing Asian American Health Research and Health Equity
   Carolyn Rubin EdD, MA, Chau Trinh-Shevrin, DrPH, Vivien Wu, and Giles Li

2. Together Strengthening the Health of Chinatown: Community Assessment of Freeway Exposure and Health (CAFEH)
   Doug Brugge, PhD, MS

3. Public Health and Health Disparities Career Discussion
   Aviva Must, PhD, Chau Trinh-Shevrin, DrPH, and Carolyn Rubin EdD, MA

   Alice Rushforth, PhD, Carolyn Rubin, EdD, MA, Harris Berman, MD, Mei-Hua Fu, MS, MEd

5. Using Data to Advance Health in Chinatown
   Giles Li, Susan Koch-Weser, ScM, ScD, Mengxi Du, Kaiyan Jew, Yuao Liu, Tina Wang, Meng Zhang

6. Results of a Healthy Chinatown Needs Assessment: Healthy Eating and Active Living Among Pre-School Aged Children in Chinatown Early Education Programs
   Virginia R. Chomitz, PhD, MS, Bernadette Davidson

7. ADAPT and Chinese Church Head Start: How a Head Start Center Benefits From Academic-Community Collaboration
   Mei-Hua Fu, MS, MEd, Merieka Torrico, MPH, Moussa Cisse, Risha DeLeon

8. Increasing Utilization of Preventative Care in Asian American Women in MA
   Lisa Gualtieri, PhD, ScM, Tam H. Nguyen, PhD, MSN/MPH, RN

9. Community Engagement to Promote Healthy Aging for Older Adults in Chinatown
   Kieran Reid, PhD, MPH

BI
Biomedical Informatics

Course description: How can you best capture the data for your study? Which electronic data capture software is best for you? Tufts CTSI’s biomedical informatics seminars review leading software applications and describe their potential for streamlining clinical trials, patient registries, surveys, and more.

1. Using REDCap™ to Build a Database or Survey
   Karen M. Switkowski, MS, MPH
CER
Comparative Effectiveness Research (CER) Survey Course

Course description: This groundbreaking course tackles one of the most opportune and relevant topics in medicine: Comparative Effectiveness Research (CER). Nationally renowned CER experts describe the current state of CER, define CER tools, and explain state-of-the-art CER methodologies in a series of 15 captivating 2-hour lectures. Each lecture has been professionally videotaped and combined with slides and other learning materials to provide interactive presentations.

1. Unit 1: Introduction
   Part 1: Comparative Effectiveness Research: Recent History and Role in Healthcare Reform
   Harry P. Selker, MD, MSPH
   Peter J. Neumann, ScD
   Part 2: Rationale for CER

2. Introduction: A Review of Evidence-Based Medicine (EBM) and a Framework for Understanding the CER Agenda
   Thomas W. Concannon, PhD

3. Unit 2: Evidence Generation
   Part 1: Comparative Effectiveness Trials
   David M. Kent, MD, MSc
   Thomas A. Trikalinos, MD, PhD
   Part 2: Assessing Pharmacogenetic Information in Clinical Trials

4. Personalized Medicine, Heterogeneity of Treatment Effect, and Implications for Comparative Effectiveness
   David M. Kent, MD, MSc

5. Retrospective and Observational Comparative Effectiveness Studies
   Dana Gelb Safran, ScD
   Peter K. Lindenauer MD, MSc

6. Unit 3: Evidence Synthesis
   Systematic Review and Meta-Analysis
   Joseph Lau, MD
   Christopher H. Schmid, PhD

7. Unit 4: Evidence Integration
   Decision Modeling – Cost-Effectiveness
   Stephen G. Pauker, MD
   John B. Wong, MD
   Decision Modeling – Simulations
   Joshua T. Cohen, PhD

8. Unit 5: Use of Evidence in Decision Making
   Community Engagement and Input into CER
   Laurel K. Leslie, MD, MPH

9. Clinical Practice Guidelines
   Katrin Uhlig, MD, MS

10. Clinical Effectiveness Trials and Predictive Instruments as Decision Support for Implementing CER
    Harry P. Selker, MD, MSPH
12. Drug Development in the CER Era  
   Kenneth I. Kaitin, PhD

13. Using Comparative Effectiveness Research to Reach Employers and Employees  
   Debra J. Lerner, MS, PhD

14. Economic and Policy Implications of CER  
   Christopher P. Tompkins, PhD

15. Unit 6: Future Directions in CER  
   Part 1: The IOM 100 Priorities and AHRQ 14 Priority Conditions and Populations  
   Part 2: The USPSTF Breast Cancer Screening Guidelines (Mammography) and CER: A Panel Discussion

CERS
Comparative Effectiveness Research (CER) Seminars

Course description: The comparative effectiveness research (CER) seminars add to the content already in the CER Survey Course. They use research examples to explain each step of the Translational Spectrum of Comparative Effectiveness Research at Tufts CTSI. The seminars also discuss the current state of CER, explain state-of-the-art CER methodologies and the importance of CER.

1. Why Comparative Effectiveness Research Matters  
   John B. Wong, MD

2. Methods for Prioritizing Research in Comparative Effectiveness Research  
   Peter Neumann, ScD, Ethan M. Balk, MD, MPH, and Joshua T. Cohen, PhD

3. Demystifying Pragmatic Clinical Trials  
   Thomas W. Concannon, PhD, Harry P. Selker, MD, MSPH, Karen Freund, MD, MPH, and Robin Ruthazer, MPH

4. Systematic Review and Meta-Analysis of Diagnostic Test Studies  
   Ethan Balk, MD, MPH and Norma Terrin, PhD

5. Making Research Useful: Dissemination and Application of Findings from Comparative Effectiveness Research  
   Dominic Hodgkin, PhD

6. Feedback & Assessment: An Essential Ingredient in Comparative Effectiveness Research  
   R. Christopher Sheldrick, PhD

7. Comparative Effectiveness Research: Where are We Going?  
   John B. Wong, MD

8. 2016 Comparative Effectiveness Research Symposium: Introduction and What’s Next, CTSI 3.0  
   John Wong, MD, and Harry P. Selker, MD, MSPH

9. The HELPS-HD Trial: A Cluster Randomized Pragmatic Trial of Oral Protein Supplements by DCI and Demystifying Pragmatic Clinical Trials  
   Daniel Weiner, MD, MS and Thomas Concannon, PhD

    Thomas Concannon, PhD, and John Wong, MD
11. Improving Screening for Diabetes in Asian Americans
Susan Koch-Weser, PhD; Anastassios G. Pittas, MD, MS; and William F. Harvey, MD, MSc, FACR

CIV
Civic Life & Health Research

Course description: What is civic life, and why is it important to consider in clinical research? Understand the relevance of civic engagement to clinical and other health research at Civic Life and Health Research, a seminar by Peter Levine, PhD, Associate Dean and Lincoln Filene Professor of Citizenship and Public Affairs at Tufts University's Jonathan M. Tisch College of Civic Life. Get oriented to current research and debates about civic engagement in the US; challenge the frequent definition of civic engagement as professionals consulting stakeholders; and instead begin to see medical researchers and health professionals as citizens who should relate to other people as fellow citizens. By the end of this seminar, participants will be able to: Define the concept of civic life and related terms such as civic engagement, social capital, democratic participation, and community engagement from social science perspectives. Value civic engagement as relevant to the clinical research process. Explore differences in theory and practice depending on whether one thinks in terms of clients, patients, stakeholders, communities, publics, or citizens. Illustrate exemplary actions that investigators can take that involve civic life.

1. Introduction to Civic Life (12 min)
   Peter Levine, PhD
2. Social Capital (16 min)
   Peter Levine, PhD
3. Collective Efficacy (6 min)
   Peter Levine, PhD
4. Common Pool Resource (8 min)
   Peter Levine, PhD
5. Public Sphere (13 min)
   Peter Levine, PhD
6. Importance in Health Research (6 min)
   Peter Levine, PhD
7. Idea of "Stakeholders" (13 min)
   Peter Levine, PhD
8. Common Tools for Civic Engagement in Research (2 min)
   Peter Levine, PhD

CMI
Common Metrics Implementation

Course description: In order to maximize the CTSA Program’s impact, the National Center for Advancing Translational Sciences (NCATS) is implementing the Common Metrics Initiative, which employs a set of common metrics for use in collaborative management based on the principles of the Results-Based Accountability (RBA) framework. The videos and activities outlined below are intended to teach you about the RBA framework, the Scorecard software and
their use/application. The materials are broken up into units that correspond with each training session and should be followed sequentially.

1. Tufts CTSI Common Metrics Implementation Session One Pre-Work
2. Tufts CTSI Common Metrics Implementation Session Two Pre-Work
3. Tufts CTSI Common Metrics Implementation Session Three Pre-Work
4. Tufts CTSI Common Metrics Implementation Session Three Post-Work

CR
Clinical Research

Course description: How is research designed, conducted, evaluated, and applied to patient care? The Clinical Research seminar series provides a comprehensive overview of clinical research methodology, from writing a research question to publishing findings. No matter what your experience, if you are interested in learning about the basic principles of research, you are encouraged to view these exciting one-hour sessions.

1. Clinical & Translational Research: An Overview and Basic Principles   
   Karen M. Freund, MD, MPH
2. Developing and Writing Research Questions, Aims & Hypotheses   
   Jonathan M. Davis, MD
3. Experimental & Observational Study Designs   
   Daniel E. Weiner, MD, MS
4. Research Using Existing Data   
   Tara Lagu, MD, MPH & Mihaela Stefan, MD, MS
5. Calculating Sample Size and Power   
   Farzad Noubary, PhD
6. Research Ethics of Clinical Investigation   
   Susan K. Parsons, MD, MRP
7. Why Studies Fail: Bias and Confounding   
   Jessica Paulus, ScD
8. Developing a Research Study Protocol   
   Tammy Scott, PhD
9. IRB and Regulatory   
   Ashley Hicks, CIP
10. Data Analysis   
    Lori Lyn Price, MAS
11. Evaluating Medical Journal Articles   
    Paul Visintainer, PhD
12. Research Data Management   
    Brian Wilson, BSc
13. Managing Multicenter Clinical Trials   
    Patricia Sheehan, RN, MS, MPH
DMRC
Developing and Managing Your Research Career

Course description: New and seasoned researchers will benefit from this outstanding professional development series that covers diverse topics such as team science, mentoring, submitting a manuscript, managing a research laboratory, and getting grant funding. These seminars are co-sponsored by the Tufts University-wide Committee for Teaching and Faculty Development.

1. **The Team Science Balancing Act: Independent Research vs. Collaborations**
   Karen M. Freund, MD, MPH

2. **Defining the Mentoring Relationship**
   Karen M. Freund, MD, MPH, Susan K. Parsons, MD, MRP, Diana W. Bianchi, MD, Michael J. Kelly, MD, MPH, and Jill L. Maron, MD, MPH.

3. **What Editors Look For in a Manuscript**
   Karen M. Freund, MD, MPH, Andrew S. Levey, MD, and Nijsje Dorman, PhD

4. **Managing a Research Laboratory**
   Karen M. Freund, MD, MPH, Sarah L. Booth, PhD, and Meghan Faherty, MS, MPH

5. **Time Management Tools and Strategies**
   Karen M. Freund, MD, MPH, Claire Weigand, and Linden T. Hu, MD

6. **Mutual Mentoring**
   Karen M. Freund, MD, MPH and Donna Qualters, PhD, MEd

7. **How to Conduct Research as a Busy Clinician**
   Karen M. Freund, MD, MPH, Eric Smith, MD, and Laura K. Snydman, MD

8. **Building a Culture of Feedback: Giving and Receiving Constructive Feedback**
   Karen M. Freund, MD, MPH, and Maria Blanco, EdD

9. **How to Manage a Large Clinical Trial**
   Susan K. Parsons, MD, MRP, Christine Wanke, MD, and Kim Dong MS, RD

10. **How to Get a K Award**
    Karen M. Freund, MD, MPH, Tara Lagu, MD, MPH, Mihaela S. Stefan, MD, and Peter K. Lindenauer, MD, MsC

11. **What To Do When You Receive Your First Grant**
    Karen M. Freund, MD, MPH, Andrew M. Hoffman, DVM, DVSc, and Robert M. Blanton, MD, MA

12. **How to Work with Industry: Insights from Experts**
    Karen M. Freund, MD, MPH, Gillian Black-Noller, MD, Jeffrey B. Blumberg, PhD, FASN, FACN, CNS, John Cosmopoulos, MSc, MBA, CLP, and Daniel G. Jay, PhD

13. **Translational Research: It's About Time**
    Karen M. Freund, MD, MPH and Harry P. Selker, MD, MPH
GCP
Good Clinical Practice for Social and Behavioral Research

Course description: The following e-Learning Course, Best Practices in Social and Behavioral Research Course, provides researchers with an overview of Good Clinical Practice (GCP) principles specific to social and behavioral clinical trials. This interactive learning resource is available courtesy of the National Center for Advancing Translational Sciences (NCATS) and the Clinical and Translational Science Awards (CTSA) Consortium. There are 9 modules in total, and each module takes approximately 25 minutes to complete. Completing the entire course is expected to take 3-4 hours. Please note that this is a self-learning tool, and may not satisfy GCP training requirements at your institution. Please check your institutional GCP training policy before completing any GCP coursework to ensure you are in compliance with NIH requirements. To receive a certificate of completion, check with your IRB office about taking this course through the “Collaborative Institutional Training Initiative (CITI) Program.

Module 1 – Introduction
Module 2 - Research Protocol
Module 3 - Recruitment and Retention
Module 4 - Informed Consent Communication
Module 5 - Confidentiality and Privacy
Module 6 - Participant Safety and AE Reporting
Module 7 - Quality Control and Assurance
Module 8 - Research Misconduct
Module 9 – Conclusion & Wrap Up

GW
Grant Writing

Course description: Looking for grant writing advice? Need help strengthening your proposals? Investigators in the process of writing a grant proposal should not miss Tufts CTSI’s grant writing seminars. With small class sizes and targeted discussions, these workshops teach participants how to attract reviewers to the significance, innovation, and approach of their studies, right from the very first page.

1. Grant Writing Strategies for Successful Research Proposals
   Amy Gantt, MA
2. Strategies for Developing a Team Science Grant Proposal
   Amy Gantt, MA
3. Writing Clearly and Concisely
   Amy Gantt, MA
HL
Health Literacy

Course description: The Tufts Medical Center Floating Hospital for Children partnered with Tufts CTSI and the American Board of Pediatrics (ABP) to implement a department-wide quality improvement (QI) project targeting residents, fellows, and faculty called “HEALERS” (Harnessing Efforts to Address Health Literacy and Enhance Relationships and Service). The project built off of materials developed by the Agency for Healthcare Quality and Research (AHRQ) and the ABP. The presentations and associated materials review what we know about Health Literacy, the purpose and operationalization of the HEALERS project, overviews of the two change strategies implemented, and a summary of HEALERS as a QI project.

1. **Addressing Health Literacy and Improving Healthcare Quality**  
   Sabrina Kurtz-Rossi, MEd and Laurel Leslie, MD, MPH

2. **Introduction to the HEALERS Project**  
   Laurel Leslie, MD, MPH, Priya Garg, MD, and Supriya Shah, BA

3. **HEALERS Change Strategy 1: Encouraging Questions**  
   Laurel Leslie, MD, MPH, Priya Garg, MD, and Supriya Shah, BA

4. **HEALERS Change Strategy 2: Teach Back**  
   Laurel Leslie, MD, MPH, Priya Garg, MD, and Supriya Shah, BA

5. **HEALERS: Putting it All Together**  
   Laurel Leslie, MD, MPH

6. **Maintenance of Certification (MOC) Overview**  
   Laurel Leslie, MD, MPH and Virginia Moyer, MD, MPH

MM
Survey, Qualitative, and Mixed Methods

Course description: What are the differences between qualitative and quantitative research, how do you decide which to use in your study? The seminars in our Survey, Qualitative, and Mixed Methods series define each research approach, explain how different methods are used in clinical and basic research, show how to design a study using qualitative and quantitative methods, and demonstrate how mixed methods can be successfully applied.

1. **An Introduction to Mixed Methods**  
   Tom Mackie, MA, PhD

2. **Mixed Methods Approaches for Health Services Research: An Introduction**  
   Justeen Hyde, PhD and Tom Mackie, MA, PhDc

3. **The Qualitative Research Process: Study Designs for Health Services Research**  
   Justeen Hyde, PhD and Tom Mackie, MA, PhDc

4. **Designing a Mixed Methods Study**  
   Justeen Hyde, PhD and Tom Mackie, MA, PhDc

5. **Applying Mixed Methods Effectively**  
   Justeen Hyde, PhD and Tom Mackie, MA, PhDc

6. **An Overview of Survey Design**  
   Susan Koch-Weser, ScD

7. **Item Response Theory**  
   Barbara Gandek, PhD
NIH
NIH Biosketch, Policy and Guidance Changes

Course description: How do you implement the recent NIH/ARHQ policy and guidance changes into your grant application? Resources such as My Bibliography and SciENcv can help you to properly format your biosketch and ensure that it is compliant with NIH policies. Experts will discuss the changes to rigor and transparency in research, inclusion reporting, data safety monitoring, vertebrate animals, definition of child, research training, appendices, biosketch clarifications, font requirements and post-award changes.

1. Navigating the New 2015 NIH Biosketch Format
   Tyler Manoukian, BA, Busra Ozturk, MEd, and Laura Schmidt, PhD

2. Planning for NIH and AHRQ Grant Application Changes
   Zoya Davis-Hamilton, EdD, CRA, Amy Gantt, MA, Kathleen Benoit, CRA and Debbie Slater, MHA

3. Uniform Guidance (UG) Implementation
   Zoya Davis-Hamilton, EdD, CRA and Joyce Ferland, BS, MBA

OH
One Health

Course description: One Health is an integrative, multi-disciplinary effort to optimize health for people, animals, and the shared environment at the local, national, and global level. The CTSI One Health program seeks to harness the synergies of diseases shared by people and animals, as well as the benefits of human-animal interactions, to advance collaborative and interdisciplinary solutions for important medical issues. This united approach will translate research into practice more effectively to optimize the health and well-being of animals, humans, and the environment. One Health incorporates expertise from diverse fields, such as human and veterinary medicine, environmental and biological sciences, engineering, public health, political science, urban planning, sociology, and statistical modeling.

1. One Health Overview
   Deborah T. Kochevar, DVM, PhD, DACVCP

2. Natural Animal Models: Beyond Rats and Mice
   Andrew Hoffman, DVM, DVSc, DACVIM, Elizabeth McNiel, DVM, PhD, DACVIM, DACVR, and Nicholas Frank, DVM, PhD

3. Healthy Pets Helping People: the Healthcare Community’s Role in Safe and Effective Animal-Assisted Therapy
   Deborah Linder, DVM, DACVN and Megan Mueller, MA, PhD

4. Zoonotic Diseases: Learning from Yesterday, Planning for Tomorrow
   Sam Telford, III, Ms, SD and Felicia Nutter, DVM, PhD

5. Environmental Aspects of One Health
   Christine L. Rioux, PhD, MS, and Antje M. F. Danielson, PhD

6. What the CTSI One Health Program Offers
   Lisa Freeman, DVM, PhD, DACVN
7. **Zoobiquity Boston Keynote Session 1: Personalizing Cancer Care for Pets and People**
   Bruce Chabner, MD

8. **Zoobiquity Boston Session 2: Obesity in a 12-Year-Old Female Domestic Shorthair Cat and a 64-Year-Old Female Pharmacist**
   Lisa Freeman, DVM, PhD, and Caroline Apovian, MD

9. **Zoobiquity Boston Session 3: Anterior Cruciate Ligament Injury in a 3-Year-Old Labrador Retriever and a 21-Year-Old Competitive Skier**
   Randy Boudrieau, DVM, and Paul Weitzel, MD

    Nicholas Dodman, BVMS, and Jean Frazier, MD

11. **Zoobiquity Boston Session 5: Peripheral T-Cell Lymphoma in a 10-Year-Old Boxer Dog and a 56-Year-Old Nurse**
    Kristine Burgess, DVM, and Andreas Klein, MD

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**PACE**

**PACE Symposium: Using Group Data to Treat Individuals**

*Course description:* What is Patient-Centered Outcomes Research and how can we improve our methods of conducting it? This Accelerating Patient-Centered Outcomes Research and Methodological Research seminar provides an overview of the Patient-Centered Outcomes Research Institute (PCORI), and the appropriate methods to implement when conducting patient-centered research.

1. **Accelerating Patient-Centered Outcomes Research and Methodological Research**
   Jessica Paulus, ScD, Harry Selker, MD, MSPH and Emily Evans, PhD, MPH

2. **Introduction to Heterogeneity of Treatment Effect (HTE)**
   David Kent, MD, MSc

3. **Risk and Treatment Effects**
   Peter Rothwell, MD, PhD

4. **Genetic Prediction of Common Diseases**
   A. Cecile J.W. Janssens, PhD

5. **Risk-Based Heterogeneity of Treatment Effect in 30 Large RCTs**
   David Kent, MD, MSc

6. **Person-Level THE**
   Issa J. Dahabreh, MD, MS

7. **Discussion on Heterogeneity Treatment Effects**
   Jessica Paulus, ScD, Ravi Varadhan, PhD, Douglas Altman, DSc, A. Cecile J.W. Janssens, PhD, and David Kent, MD, MSc

8. **A Proposed Guideline for Reporting HTE in Large Randomized Clinical Trials**
   Rod Hayward, MD

9. **Panel Discussion**
   Ewout Steyerberg, PhD, Douglas Altman, DSc, Robert Golub, MD, Rod Hayward, MD, David Kent, MD, MSc, and Walter Kernan, MD
Quality Improvement in Healthcare

Course description: This course will provide you an overview of the Quality Improvement (QI) methods and tools, and how can you apply them to your research? These seminars begin with an overview of QI, developing your aims statement, QI tools, and then various examples of how you implement small changes, how you measure change, and how you lead change.

1. What is Quality Improvement in Healthcare
   Denise Daudelin, RN, MPH
2. The Power of Quality Improvement
   Laurel K. Leslie, MD, MPH and Carmina Erdei, MD
3. Quality Improvement Tools: An Overview
   Priya Garg, MD
4. Aims Statements: Case Examples
   Denise Daudelin, RN, MPH
5. Measuring Change Strategies: Decreasing Air Leaks and Testing with PDSA Cycles
   Laurel K. Leslie, MD, MPH and Patoula Panagos, MD
6. Implementing Change Strategies: Patient Falls
   Denise Daudelin, RN, MPH and Tricia Ide, RN, MS
7. Using Run Charts: Readmissions
   Laurel K. Leslie, MD, MPH and Alexander Pavoll, MPH
8. Statistical Process Control
   Munish Gupta, MD, MMSc
9. Team Charters, Team Dynamics, and Leading Change
   Laurel K. Leslie, MD, MPH and Denise Daudelin, RN, MPH
10. Creating High Reliability for Organizational Improvement
    Evan Benjamin, MD, FACP and Stephanie Calcasola, MSN, RN-BC

Regulatory Affairs

Course description: Why do you need to provide informed consent to clinical research subjects? What are the Principal Investigator’s responsibilities for a clinical research study? What are successful strategies in working with the Institutional Review Board (IRB)? Regulatory affairs seminars provide information on how clinical research studies need to ensure subject safety and welfare while complying with federal, state and institutional regulations.

1. Principal Investigator IRB Responsibilities
   Andreas K. Klein, MD
2. Working with the IRB: Common Myths and Successful Strategies
   Ashley D. Hicks, CIP
3. Best Practices in Clinical Trials
   David R. Snydman, MD, FACP, FIDSA
4. Navigating Contracts and Agreements
   Frederick M. Frankhauser, JD, MBA, RPh
5. **Collaborative Institutional Review Board (IRB) Agreements**  
   Andreas K. Klein, MD, Gordon S. Huggins, MD, Ashley D. Hicks, CIP, and Jonathan M. Davis, MD

6. **Master Contracts**  
   Paul Murphy, JD, MPA

7. **Data Safety and Monitoring Boards: A Brief Overview**  
   Tamsin A. Knox, MD, MPH

8. **What You Need to Know About DSMBs (But Were Afraid to Ask)**  
   Tamsin A. Knox, MD, MPH

9. **Data Safety and Monitoring Board Information for IRB Members**  
   Tamsin A. Knox, MD, MPH

10. **The Challenge of Informed Consent: A Proposal**  
    Jonathan M. Davis, MD

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**RC**  
**Research Coordinator Education**

*Course description:* In this series for study coordinators, research assistants, and other members of the research team: the Research Coordinator Education Program. This series will outline the roles and responsibilities of the research team throughout a research project.

1. **Clinical Research Lifecycle**  
   Tamsin Knox, MD, MPH

2. **Clinical Trial Lifecycle**  
   Veronika Testa, BSN, RN, CCRC and Douglas Reichgott

3. **Clinical Research Billing and Medicare Coverage Analysis**  
   Douglas Reichgott

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**RDDA**  
**Research Design and Data Analysis**

*Course description:* How many subjects do you need for your study? What’s a P-value? How do you handle missing data? These are just some of the questions answered in our statistical, epidemiological, and research-oriented seminar series. Experienced instructors from Tufts CTSI’s Research Design Center/Biostatistics Research Center (RDC/BRC) will help you to avoid common pitfalls and learn to build a successful research career.

1. **Introduction to Study Design**  
   Jessica Paulus, ScD

2. **Formulating Research Questions, Hypotheses and Objectives**  
   Kahsi A. Smith, PhD

3. **Concepts of Hypothesis Testing**  
   Lori Lyn Price, MAS

4. **How Many Subjects Do I Need for My Study?**  
   Jessica K. Paulus, ScD

5. **Developing a Study Protocol**  
   Tammy Scott, PhD
6. **Pitfalls in Statistical Analysis**  
   Lori Lyn Price, MAS  
7. **Bias and Confounding in Clinical Research**  
   Jessica Paulus, ScD  
8. **Linear and Logistic Regression**  
   Lori Lyn Price, MAS  
9. **Modeling Time-to-Event Outcomes**  
   Robin Ruthazer, MPH  
10. **Which Statistical Test Should I Use?**  
    Lori Lyn Price, MAS

**RPI**  
Research Process Improvement

*Course description:* Process improvement approaches (e.g. LEAN, Six Sigma, Model for Improvement) have been employed for years in industry to improve the quality, efficiency, and effectiveness of processes. What are the current trends, methods, tools and resources in process improvement and how can you apply them to your research? These seminars begin with an overview of process improvement, take a close look at tools used in process and quality improvement and how to use them, and feature a discussion with other researchers on how they addressed challenges in their research applying these approaches.

1. **Deconstructing Quality Improvement and Applying it to Research**  
   Laurel Leslie, MD, MPH and Denise Daudelin, RN, MPH  
2. **Using QI Tools for Improving Research Efficiency and Reliability**  
   Laurel Leslie, MD, MPH and Denise Daudelin, RN, MPH  
3. **Using QI Tools for Managing Research Teams and Participant Recruitment and Retention**  
   Laurel Leslie, MD, MPH and Denise Daudelin, RN, MPH

**RRP**  
Research Recruitment and Participation

*Course description:* Why do people participate in clinical studies and trials? What are the challenges to recruiting human and animal research participants, and how can we overcome them? Find out at though this series of lectures delivered at Tufts CTSI’s Translational Research Day: Innovations in Clinical Trial Participant Engagement.

1. **Introduction, The mPower App and Using Technology for Participant Engagement in Clinical Trials**  
   Harry Selker, MD, MSPH, Karl Kieburtz, MD, MPH  
2. **Participation in Clinical Research: Motivations and Perspectives**  
   Julia Farides-Mitchell, MA  
3. **Companion Animal Studies: Participant Engagement**  
   Andrew M. Hoffman, DVM, DVSc  
4. **Challenges in Consenting Pregnant Women, Children and Neonates**  
   Jonathan Davis, MD
5. **Participant Engagement Panel**  
   Julia Farides-Mitchell, MA, Andrew M. Hoffman, DVM, DVSc, Jonathan Davis, MD

6. **Using Social Media for Participant Engagement in Clinical Trials: A Pilot Study**  
   Laura Blaisdell MD, MPH, FAAP

7. **Recruitment in Social Behavioral Research (Dear Abby and More)**  
   Debra Lerner, MS, PhD

8. **Using and Expert Panel to Randomize Patients in a Cervical Spondolytic Myelopathy Clinical Trial**  
   Zoher Ghogawala, MD

9. **Innovative Recruitment Strategies Panel**  
   Laura Blaisdell MD, MPH, FAAP, Debra Lerner, MS, PhD, Zoher Ghogawala, MD

10. **Using Research Process Improvement to Solve Recruitment Challenges**  
    Denise Daudelin, RN, MPH, Donato Rivas, PhD

**SE**

**Stakeholder and Community Engagement**

*Course description:* Engagement of stakeholders is increasingly called for in federal and foundation-funded research. Who are these community stakeholders, and why is it important to involve them in your study? We will define patient, stakeholder, and community engagement in research, discuss national trends among funding agencies with respect to engagement of the public in research, describe the Tufts typology of stakeholder types (the 7 Ps), and provide an approach for mapping your research needs to different models of engagement of the public in research.

1. **Stakeholder and Community Engagement: Why It's Important**  
   Laurel K. Leslie, MD, MPH

2. **Stakeholder and Community Engagement: Lessons Learned**  
   Laurel K. Leslie, MD, MPH and Carolyn Rubin EdD, MA

3. **Methods of Stakeholder Engagement**  
   Carolyn Leung Rubin, EdD, MA and Laurel K. Leslie, MD, MPH

4. **Community Engagement to Improve Asian Health**  
   Carolyn Rubin EdD, MA and Mei-Hua Fu, MS, MEd

5. **Stakeholder Engagement in Patient-Centered Comparative Effectiveness Research**  
   Thomas Concannon, PhD

6. **Preparing for Patient-Centered and Stakeholder-Engaged Research**  
   Thomas Concannon, PhD

7. **Civic Life and Health Research**  
   Thomas Concannon, PhD and Peter Levine, PhD

8. **Engaging Stakeholders in Community-Based Participatory Research Partnerships**  
   Thomas Concannon, PhD and Carolyn Rubin, EdD, MA

**TRD**

**Translational Research Day 2017: Sensors, Devices, and Biomarkers in Medicine**

*Course description:* How can sensor, device, and biomarker data improve health, prevent and detect disease at an earlier stage, and personalize interventions? Find out at Tufts CTSI's
Translational Research Day 2017: Sensors, Devices, and Biomarkers in Medicine. The learning objectives for Translational Research Day include that viewers of the day’s talks will be able to: Recognize the different classifications of biomarkers and their potential in detecting early-stage disease and for personalizing interventions; Illustrate diverse approaches to advancing the capabilities of sensors and medical devices and their practical applications in improving health; Describe potential translational roadblocks in developing, testing, and using sensor- and device-based health prevention, detection, management, and intervention strategies; Identify Tufts CTSI resources and services that support team-based translational science.

1. Improving the Assessment of Functional Change in CNS Clinical Trials (15 min)
   Josh Cosman, PhD
2. Smart Mechanical Support Devices for Cardiac Care (11 min)
   Navin Kapur, MD
3. In Vivo Nanosensors and Imaging Technologies (12 min)
   Heather Clark, PhD
4. Improving Behavioral Measurements from Mobile Devices (15 min)
   Stephen S. Intille, PhD
5. Embedded Functioning of Nanoscale Sensors in Hybrid Tissues (13 min)
   Brian Timko
6. Harmonizing Biomarker Terminology: NIH-FDA BEST (Biomarkers, EndpointS, and other Tools) (22 min)
   Christopher Leptak, MD, PhD
7. A Metatranscriptomic Approach to Salivary Biomarker Discovery in the Premature Newborn (13 min)
   Jill L. Maron, MD, MPH
8. Collaborative Data Science in Health Care (14 min)
   Leo Anthony Celi, MD, Msc, PhD
   Kumaran Kolandaivelu, MD, PhD
10. Lessons Learned from the Front Lines (24 min)
    Rami Tzafriri, PhD and Michael Naimark, MS
11. Case Study Panel Discussion (12 min)
    Rami Tzafriri, PhD; Michael Naimark, MS; Kumaran Kolandaivelu, MD, PhD
12. Funding Opportunities - Tufts CTSI Symposium Plus (12 min)
    Graham Jones, PhD; Alysse Wurcel, MD, MS; John Leong, MD, PhD

TSS
Translational Science Seminars

Course description: This series will explore Clinical and Translational Science across its full spectrum, specific contributions of different parts of the spectrum in the overall goal of improving health, and understand topic-specific examples that one can draw upon to exemplify translational research. The seminar series will highlight current research being done cross-collaboratively between Tufts CTSI and across the Tufts campus, its community and affiliates.

1. Exploiting the Molecular Signatures of Disease: Case Studies in Bench-to-Bedside Research
   Graham B. Jones, PhD, DSc
2. Bile Acids and Spore Germination: A Novel Approach to Blocking Clostridium Difficile Infection
   Abraham L. Sonenshein, PhD and Yoav Golan, MD, MS
3. Translating ‘Natural’ Experiments into Clinical Research: Using Administrative Data for CER
   Daniel E. Weiner, MD, MS
4. Spontaneous Animal Models in Translational Research: From Cardiac Cachexia to Obesity
   Lisa Freeman, DVM, PhD, DACVN

TTIC
Technology Transfer and Industry Collaboration

Course description: As part of our continuing efforts to support research, innovation, translation, entrepreneurship and technology transfer within the Tufts community, Tufts Technology Transfer and Industry Collaboration (TTIC) has organized a seminar series of lectures on intellectual property management, technology transfer, start-up company formation and other relevant subjects. The topics are planned for experience at all levels and provide an overview of the role of university technology tran

1. Introduction to Technology Transfer
   Erik Halvorsen, PhD, MBA
2. An Introduction Into Material Transfer Agreements
   Lee Tien, PhD
3. The Invention Disclosure Process
   Colm Lawler, PhD
4. An Introduction Into License and Option Agreements
   Martin Son, PhD
5. How to Start a Company Based on a University Technology
   Paul Hartung, MS
6. Negotiation Techniques
   Erik Halvorsen, PhD, MBA
7. Effective Marketing Strategies
   Jennifer Tsai, MBA
8. The Role of Technology Development for Advancing Academic Discoveries
   John Cosmopoulos, MSc, MBA
9. Business Communication
   Erik Halvorsen, PhD, MBA
    Erika Bechtold, PhD