Doctoral Students and Alumnus of the USC Regulatory Science Program. Some thesis titles are working titles.

Name	Thesis/Dissertation Title
Lilit Aladadyan	Investigating Parental Attitudes Toward Food Advertising Directed at Children: Factors Influencing Household Good Purchases
Dr. Susan Bain	"Regulatory" Due Diligence: A Survey Investigation of Best Practices in the Medical Products Industry
Dr. Tony Chan	Implementation of Risk Management in Medical Device Companies: A Survey Analysis of Current Practices
Terry Church	Biobank Topography and the Contours of Community Governance: Mapping the Moral and Regulatory Landscape
Alicia Colarossi	Risk Evaluation Mitigation (REMS) in the US
Grant Dagliyan	Development of New Radiopharmaceuticals for Positron Emission Tomography: Challenges and Opportunities Perceived by Imaging Thought Leaders
Kobby Dankwah	Considerations for Social Media Promotion of Regulated Products
Dimapindan, Patrick	An Integrated Model and Framework for Evaluation of Customer Friendly Delivery: A Study of Electronic Systems at FDA's Los Angeles Import Operations
Clare Elser	Validation Master Planning Impacts on Compliance Profile
Aimee Greco	Regulatory Dissonance with the European CE Mark Process and US FDA Regulations for Class II Medical Devices
Grant Griffin	The Russian Federation as an Emerging Market for Pharmaceutical and Medical Device Clinical Trials: A Survey of Industry Views
Isara Isarowong	Effectiveness of Current Anti-Counterfeit Measures in Pharmaceutical Products
Dr. Michael Jamieson	The Role of Universities in the Commercialization of Medical Products: A Survey of Industry Views
Martha Kamrow	US Perception of Changing Regulations in Emerging Markets for Class II Medical Devices
Bill Leitner	The Impact of the FDA on Advisory Panel Outcomes: A Survey of Industry Views
David Locke	Educational Needs of Regulatory Professionals: A Survey of Industry Professionals with Healthcare Product Regulatory Experience
Dr. Duane Mauzey	Current Practices in Pharmaceutical Container Closure Development
Haven (Richard) McCall	Risk Management Approaches and Standards Used in US Hospitals
Cesar Medina	Integrating Patient-Focused Benefit-Risk Assessments in the U.S. Drug Approval Decisions: A Case Study on Risk Tolerance of Drugs Used to Treat Autism in Children

Caroline Mosessian,	Health Technology Assessment: How do Hospital Executives and Physicians Decide and Value Implant Selection
Vada Perkins	To Determine the Cause of Underreported Adverse Drug Events/Reactions by Patients and Healthcare Professionals in the US
JoAnn Pfeiffer	The Role of the Academic Investigator in the Review and Negotiation of the Clinical Trial Agreement: A Survey of Investigators' Knowledge, Engagement and Motivation
Susan Pusek	Strengthening the Regulatory Workforce: Defining Optimal Training in Regulatory Science for Participants in NIH-funded Institutional K-12 Programs
Dr. Valerie Ramsey	Clinical Trials Driven by Investigator-Sponsors: GCP Compliance With or Without Previous Industry Sponsorship
Ali Rejaei	Regulatory Challenges Associated with Botanical Products for Therapeutic use, A survey of Industry
Catherine Sheehan	International Harmonization of Pharmacopeial Standards: A Survey Investigation of the Pharmacopeial Discussion Group's (PDG) Key Challenges
Dr. Taranjit Singh	Software Risk Management: An Exploration of Software Life Cycle Methodologies, Best Practices and Tools for their Application to Medical Device Software Risk Management
Nancy Smerkanich	A Comparison of Qualitative vs. Quantitative Benefits/Risks Framework for New Drugs
Alexa Smith	Affects of Recent EU Color Legislation and Consumer Demand for Natural Colors on the Pharmaceutical and Dietary Supplement Industries
Dr. Martin Solberg	A Survey of Transparency in Three Asian Regulatory Agencies Responsible for Medical Products
Chin-Wei Soo	Regulatory Challenges Governing Patient Access to Humanitarian Use Devices
Neal Storm	Regulatory Dissonance in the Global Development of Drug Therapies: A Case Study of Drug Development in Postmenopausal Osteoporosis
Simone Turnbull	Legalizing Cosmeceuticals: Proposing a Policy Framework for how Cosmeceuticals should be Regulated in the United States
Mauricio Umana	Stem Cell Therapy: Critical Assessment of FDA Current Regulation and Guidance Documents
Loren Wagner	Regulatory Obstacles to Pharmaceutical Research and Development Productivity
Dr. Ellen Whalen	The Impact of Incomplete Monographs on the OTC Drug Industry: A Survey Investigation of Industry Views