

MARCH 28, 2024



SPA CERTIFICATION PROGRAM

Proposed Syllabus

PRESENTED BY: ALINA GRAYS

ASSISTANT DIRECTOR, SPA EDUCATION & BUSINESS PROCESS

FY25 Session Overview:

- **Application process:** Begin advertising Session 1 (starting in Sept. 2024) in Research Roundup and SPA Weekly Updates for 2 months (starting mid-April). Applications due by mid-July. Application review and acceptance notifications by mid-August.
- **Class size:** A maximum of 18 students will be admitted to the program each fiscal year.
- **What do participants bring to each session?** Laptop, calculator, paper/pen
- **Contract:** commitment of 12 months for all participants, with provision for if they decide the program is not for them at some point along the way.
- **Session Length:** 8:30 AM – 12:30 PM, once monthly
- **Classroom Location:** Trinity Towers Classroom (all in-person sessions)
- **Metrics:** Plan to track attendance, passage of Session Tests and will request feedback for each session (post-session evaluation/survey). Will send surveys to Supervisors and colleague of their choosing (Chair, PI, DA, etc.) at conclusion of 12-months for feedback and if they saw marked improvement/growth.
- **Sessions Tests:** Combination of multiple choice, essay, short-answer and problem solving (as indicated by curriculum for each session).
- **SPA Certification:** Certificates given at end of 12-month program for those who completed and have a cumulative passing score.

Session Schedule:

Basic Science: 8-month program (starting Sept. 2024)

- **September 2024- Session 1**
 - Pre-requisite video to include:
 - Introduction to Research Administration
 - Define research administration
 - Job description of the research administrator
 - What is the research enterprise?
 - History of research administration
 - The future of research administration
 - Infrastructure for research administration
 - Code of Federal Regulations & Uniform Guidance
 - Pre-Award Session 1:
 - Introduce concept of pre-award and proposal development
 - Examine elements of a successful proposal
 - Understand proposed budget, types of budgets, and budget justification

- Q&A
- Session 1 Test

- **October 2024- Session 2**
 - Pre-Award Session 2:
 - Identify/define key personnel
 - Understand indirect costs
 - Review UTSW/SPA signature authority
 - Navigate eGrants

 - Q&A
 - Session 2 Test

- **November 2024- Session 3**
 - Pre-Award Session 3:
 - Funding Agency Overview
 - National Institutes of Health (NIH)
 - Department of Defense (DOD)
 - Cancer Prevention and Research Institute of Texas (CPRIT)
 - Welch Foundation
 - American Heart Association (AHA)

 - Q&A
 - Session 3 Test

- **December 2024- Session 4**
 - Pre-Award Session 4:
 - Introduce non-industry agreement execution
 - Examine federal research contracts/subawards
 - Gain understanding of contract types and potential issues
 - Understand contract amendments/modifications
 - Review terms and conditions
 - Understand subrecipient monitoring and risk mitigation
 - Examine challenges/opportunities of international research

 - Q&A
 - Session 4 Test

- **January 2025- Session 5**
 - Post-Award Session 1:
 - Facilities & Administrative Costs
 - Cost Principles
 - Compliance with Regulations
 - Award Acceptance
 - Review of NOA, terms & conditions, special terms & conditions
 - Setting up a COA and receiving GNR
 - What's next?

 - Q&A
 - Session 5 Test

- **February 2025- Session 6**
 - Post-Award Session 2:

- Award Maintenance
 - Team interactions
 - Award modifications
 - Expense monitoring and cost transfers
 - Error correction and transactions
 - Unobligated balance - Planning for year-end
- Revenue & Billing
 - Bill plans overview
 - Collections and Refunds
- Q&A
- Session 6 Test
- **March 2025- Session 7**
- Post-Award Session 3:
 - Financial Reporting
 - Reporting Overview
 - Managing finances and financial reporting
 - eGrants
 - Special considerations
 - Award Closeout
 - Monthly monitoring and financial closeout
- Q&A
- Session 7 Test
- **April 2025- Session 8**
- Post-Award Session 4:
 - Hold time for any residual topics or questions, ad-hoc
- Compliance and Audits
 - Compliance Issues
 - Audits
 - Working with Internal and External Auditors
 - Salary & Wage Confirmation
 - Data Management
 - Responsible Conduct of Research
 - Conflict of Interest in Research
- Q&A
- Session 8 Test

Clinical Research: 4-month program (starting May 2025)

- **May 2025- Session 9**
 - Pre-requisite video to include:
 - Introduction to Clinical Research Administration
 - Define clinical research
 - Types of studies – Fundamentals of clinical trial design
 - Job description of the clinical research administrator
 - History of clinical research administration
 - The future of clinical research administration
 - Infrastructure for clinical research administration

- Clinical Research Overview: Lifecycle of clinical research
 - Clinical Trials: getting started
 - Informed Consent
 - Central office components and Trainings/Resources: CRS/CT Finance/HRPP/OCR
- Q&A
- Session 9 Test

- **June 2025- Session 10**
 - Industry Agreements- Types of agreements, timing of execution, etc.
 - Terms & Conditions
 - Risk Mitigation
 - Intellectual Property Considerations
 - Signature Authority
 - Q&A
 - Session 10 Test

- **July 2025- Session 11**
 - Coverage Analysis Overview
 - Congruency Review
 - Timing of execution
 - Calendaring
 - Introduce Grants/Contracts Mentorship Program & application deadlines- Jennifer Pitman
 - Q&A
 - Session 11 Test

- **August 2025- Session 12**
 - Clinical Trial Finance Overview
 - Clinical Trial Budget Development
 - Invoicing for Clinical Trial Activities (DCT)
 - ClinCard overview and best practices
 - Q&A
 - Session 12 Test