

TEMPLE UNIVERSITY SCHOOL OF PHARMACY REGULATORY AFFAIRS AND QUALITY ASSURANCE GRADUATE PROGRAM

POST MASTER'S **CERTIFICATE IN CLINICAL TRIAL MANAGEMENT**

APPLICATION

- 1. Students must formally apply to Post-Master's Certificates before registering for courses.
- 2. Mail hard copy of this form to Temple U, RA and QA Graduate Program, 425 Commerce Drive, Suite 175, Fort Washington, PA 19034.
- 3. Include photocopies of transcripts from all undergraduate and graduate colleges and universities attended.
- 4. Certificates are not automatically awarded. You must submit the Notice of Completion (available on the Certificate Link) to the RAQA Office by the stipulated deadline.



Name		
Address	Apt	
City	State	Zip
TUid (Temple MS graduates only)	_ E-mail Address	
Daytime phone		
Undergraduate School attended		
Degree Received	Year	
Graduate School attended	Year	
Signature	Date	
On a separate sheet of paper, please write a brief statement (maximum 350 words) of why you are interested in pursuing the Post-Master's Certificate in Clinical Trial Management.		
If you already received the MS in RAQA from Temple U, but did not complete one (or all) of the following four courses, you must complete them to receive the Post-Master's Certificate in Clinical Trial Management. Check which one you intend to take. (Students who did not receive the MS in RAQA must take all four plus one additional elective):		
☐ Good Clinical Practices (5536)		
☐ Clinical Trial Management (5537) or Global Clinical Drug Development (5539)		
☐ Bioethics for Pharmaceutical Professionals (5612)		
☐ Statistics for Clinical Trials (5497) OR ☐ Clinical Data ☐ Clinical Drug Safety and Pharmacovigilance (5538) OR ☐ Risk Management and Safety Signaling in Healt	OR	Clinical Trials (5547)
If you have already taken all of the above listed courses as part from the following electives as a substitute. Please indicate wh		u may select a course(s)
☐ Fundamentals of Pharmacology and Pharmacokinetics (5401)*	☐ Pharmacoepidemiology (5573)
Quality Audit (5494)*	☐ Risk Management and Safety S	Signaling in
Clinical Drug Safety and Pharmacovigilance (5538)	Healthcare Products (5578)	DI : 1 (5550
☐ Project Management for Clinical Trials (5547) ☐ Post Marketing Safety Surveillance (5571)*	☐ Regulatory and Legal Basis of ☐ Clinical Aspects of Pharmaceu	•
* These courses count towards the Post-Master's Clinical Trial N	•	

the MS in RAQA.