



Individual Courses and Bundles

Mr Ms Dr First Name _____ MI _____ Last Name _____

Advanced Degree: JD PhD PharmD MD DDS DMD SCD DVM RAC

Title _____

Company _____

Address Business _____ Suite/Apt _____
 Home

City/ State/Province _____

Mail Stop _____ Postal Code _____ Country _____

Phone (with area/country code) _____

Email Address _____

Billing Address (if different from above) Business _____ Suite/Apt _____
 Home

City/ State/Province _____

Mail Stop _____ Postal Code _____ Country _____

By registering for this event, I hereby agree that my profile will be stored with RAPS and shared with event vendors for the purpose of administering this event. I understand that I may receive emails from RAPS regarding this event and future related events and that I can opt-out at any time by contacting RAPS or updating preferences.
I also understand that I will be added to an online community on RAPS Regulatory Exchange at connect.raps.org, powered by Higher Logic for networking and related resources and announcements regarding this event. I may opt out at any time on connect.raps.org.

REGISTRATION FEES (All fees in US dollars)

CLINICAL	Member	List	Enterprise
Globalization of Clinical Research Trials and Investigations	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$328
Good Clinical Practice (GCP)	<input type="checkbox"/> \$255	<input type="checkbox"/> \$350	<input type="checkbox"/> \$229
Understanding and Managing the US Clinical Trial Process	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$418
ESSENTIALS	Member	List	Enterprise
Effective Regulatory Communication	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$328
Ethics	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$418
FDA Law and Regulation	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$418
Intermediate Medical Writing: Pharmaceuticals and Biologics	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$328
Intermediate Medical Writing: Investigational Applications	<input type="checkbox"/> \$570	<input type="checkbox"/> \$790	<input type="checkbox"/> \$513
Intermediate Medical Writing: Medical Devices	<input type="checkbox"/> \$255	<input type="checkbox"/> \$350	<input type="checkbox"/> \$229
Introduction to Regulatory Affairs in the EU	<input type="checkbox"/> \$135	<input type="checkbox"/> \$185	<input type="checkbox"/> \$121
Introduction to Regulatory Affairs in the US and Canada	<input type="checkbox"/> \$255	<input type="checkbox"/> \$350	<input type="checkbox"/> \$229
Introductory Medical Writing	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$328
Project Management for Regulatory Professionals	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$418
Regulatory Due Diligence for Product Development	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$328
Role of the Regulatory Professional	<input type="checkbox"/> \$ 25	<input type="checkbox"/> \$ 25	<input type="checkbox"/> \$22
Supplier Management	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$328
Supply Chain Controls	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$328
MEDICAL DEVICES	Member	List	Enterprise
Global Regulatory Strategy for Medical Devices	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$418
Medical Devices: Advertising and Promotion in the US	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$328
Medical Devices: Canadian Regulations	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$328
Medical Devices: China, Japan, Singapore and South Korea Regulatory Overview	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$418
Medical Devices: Compliance and Audits	<input type="checkbox"/> \$545	<input type="checkbox"/> \$745	<input type="checkbox"/> \$490
Medical Devices: Corrections, Removals and Directed Recalls	<input type="checkbox"/> \$570	<input type="checkbox"/> \$790	<input type="checkbox"/> \$513
Medical Devices: Definition and Lifecycle	<input type="checkbox"/> \$135	<input type="checkbox"/> \$185	<input type="checkbox"/> \$121
Medical Devices: EU Regulations	<input type="checkbox"/> \$545	<input type="checkbox"/> \$745	<input type="checkbox"/> \$490
Medical Devices: Postmarket Surveillance	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$418
Medical Devices: Risk Management	<input type="checkbox"/> \$465	<input type="checkbox"/> 640	<input type="checkbox"/> \$418
Medical Devices: US Regulations	<input type="checkbox"/> \$545	<input type="checkbox"/> \$745	<input type="checkbox"/> \$490
Regulation of Combination Products	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$328
Regulation of IVDs in the US and Major Markets Outside the US	<input type="checkbox"/> \$545	<input type="checkbox"/> \$745	<input type="checkbox"/> \$490



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CUSTOMER NAME

First _____ MI _____ Last _____

PHARMACEUTICALS	Member	List	Enterprise
Chemistry, Manufacturing and Controls	<input type="checkbox"/> \$545	<input type="checkbox"/> \$745	<input type="checkbox"/> \$490
Global Regulatory Strategy for Pharmaceuticals	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$418
Pharmaceuticals: Advertising and Promotion in the US	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$328
Pharmaceuticals: Canadian Regulations	<input type="checkbox"/> \$570	<input type="checkbox"/> \$790	<input type="checkbox"/> \$513
Pharmaceuticals: Compliance and Audits	<input type="checkbox"/> \$545	<input type="checkbox"/> \$745	<input type="checkbox"/> \$490
Pharmaceuticals: Definition and Lifecycle	<input type="checkbox"/> \$135	<input type="checkbox"/> \$185	<input type="checkbox"/> \$121
Pharmaceuticals: EU Regulations	<input type="checkbox"/> \$570	<input type="checkbox"/> \$790	<input type="checkbox"/> \$428
Pharmaceuticals: US Regulations	<input type="checkbox"/> \$545	<input type="checkbox"/> \$745	<input type="checkbox"/> \$409
Pharmacovigilance	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$418
Regulation of Biosimilars	<input type="checkbox"/> \$255	<input type="checkbox"/> \$350	<input type="checkbox"/> \$229
Regulation of Combination Products	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$328
Regulation of Dietary Supplements and NHPs	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$328
Regulation of Generic Drugs in the US	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$328
Regulation of US and EU Biologics	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$418
REMS and RMPs	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$328
QUALITY	Member	List	Enterprise
Good Laboratory Practice (GLP)	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$328
Good Manufacturing Practice (GMP)	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$418
Quality System Regulation (QSR)	<input type="checkbox"/> \$545	<input type="checkbox"/> \$745	<input type="checkbox"/> \$490

REGISTRATION FEES (All fees in US dollars)

BUNDLES	Member	List	Enterprise
Clinical Trial Foundations	<input type="checkbox"/> \$ 575	<input type="checkbox"/> \$790	<input type="checkbox"/> \$518
GxP	<input type="checkbox"/> \$ 865	<input type="checkbox"/> \$1292	<input type="checkbox"/> \$779
Regulatory Basics: Complete	<input type="checkbox"/> \$ 500	<input type="checkbox"/> \$ 700	<input type="checkbox"/> \$450
Regulatory Basics: US and Canada	<input type="checkbox"/> \$ 400	<input type="checkbox"/> \$ 560	<input type="checkbox"/> \$360
Regulatory Medical Writing: Complete	<input type="checkbox"/> \$1240	<input type="checkbox"/> \$1710	<input type="checkbox"/> \$1116
Regulatory Medical Writing: Package #1	<input type="checkbox"/> \$ 1050	<input type="checkbox"/> \$1440	<input type="checkbox"/> \$878

METHOD OF PAYMENT

International Wire Transfer: Fax a completed form and copy of bank wire confirmation to confirm your registration to the following (a \$30 administration fee will apply):

RAPS account #10000432281257—ABA #061000104—Swift Code SNTRUS3A to: SunTrust Bank, Richmond, VA. Must reference name of registrant. All bank charges are the responsibility of the payer.

Check # _____

Credit Card American Express MasterCard Visa
Account # _____ Exp. Date _____ Billing Postal Code _____

Name as it appears on the card _____ Signature _____

Questions? Call RAPS Solutions Center at +1 301 770 2920, ext. 200. Please see **RAPS.org** for complete registration policies and procedures.

HOW TO REGISTER

ONLINE: <https://www.raps.org/events-training/online-training-and-certificates> (credit card only)

MAIL: RAPS 5635 Fishers Lane, Suite 550, Rockville, MD 20852.

FAX: +1 301 841 7956 (credit card or wire)