



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"/> \$274
Good Clinical Practice (GCP)	<input type="checkbox"/> \$255	<input type="checkbox"/> \$350	<input type="checkbox"/> \$191
Understanding and Managing the US Clinical Trial Process	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$349
ESSENTIALS	Member	List	Enterprise
Effective Regulatory Communication	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$274
Ethics	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$349
FDA Law and Regulation	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$349
Intermediate Medical Writing: Pharmaceuticals and Biologics	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$274
Intermediate Medical Writing: Investigational Applications	<input type="checkbox"/> \$570	<input type="checkbox"/> \$790	<input type="checkbox"/> \$428
Intermediate Medical Writing: Medical Devices	<input type="checkbox"/> \$255	<input type="checkbox"/> \$350	<input type="checkbox"/> \$191
Introduction to Regulatory Affairs in the EU	<input type="checkbox"/> \$135	<input type="checkbox"/> \$185	<input type="checkbox"/> \$101
Introduction to Regulatory Affairs in the US and Canada	<input type="checkbox"/> \$255	<input type="checkbox"/> \$350	<input type="checkbox"/> \$191
Introductory Medical Writing	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$274
Project Management for Regulatory Professionals	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$349
Regulatory Due Diligence for Product Development	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$274
Role of the Regulatory Professional	<input type="checkbox"/> \$ 25	<input type="checkbox"/> \$ 25	<input type="checkbox"/> \$19
Supplier Management	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$274
Supply Chain Controls	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$274
MEDICAL DEVICES	Member	List	Enterprise
Global Regulatory Strategy for Medical Devices	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$349
Medical Devices: Advertising and Promotion in the US	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$274
Medical Devices: Canadian Regulations	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$274
Medical Devices: China, Japan, Singapore and South Korea Regulatory Overview	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$349
Medical Devices: Compliance and Audits	<input type="checkbox"/> \$545	<input type="checkbox"/> \$745	<input type="checkbox"/> \$409
Medical Devices: Corrections, Removals and Directed Recalls	<input type="checkbox"/> \$570	<input type="checkbox"/> \$790	<input type="checkbox"/> \$428
Medical Devices: Definition and Lifecycle	<input type="checkbox"/> \$135	<input type="checkbox"/> \$185	<input type="checkbox"/> \$101
Medical Devices: EU Regulations	<input type="checkbox"/> \$545	<input type="checkbox"/> \$745	<input type="checkbox"/> \$409
Medical Devices: Postmarket Surveillance	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$349
Medical Devices: Risk Management	<input type="checkbox"/> \$465	<input type="checkbox"/> 640	<input type="checkbox"/> \$349
Medical Devices: US Regulations	<input type="checkbox"/> \$545	<input type="checkbox"/> \$745	<input type="checkbox"/> \$409
Regulation of Combination Products	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$274
Regulation of IVDs for Key International Markets	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$349
Regulation of IVDs in the US	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$274



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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"/> \$409
Global Regulatory Strategy for Pharmaceuticals	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$349
Pharmaceuticals: Advertising and Promotion in the US	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$274
Pharmaceuticals: Canadian Regulations	<input type="checkbox"/> \$570	<input type="checkbox"/> \$790	<input type="checkbox"/> \$428
Pharmaceuticals: Compliance and Audits	<input type="checkbox"/> \$545	<input type="checkbox"/> \$745	<input type="checkbox"/> \$409
Pharmaceuticals: Definition and Lifecycle	<input type="checkbox"/> \$135	<input type="checkbox"/> \$185	<input type="checkbox"/> \$101
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"/> \$349
Regulation of Biosimilars	<input type="checkbox"/> \$255	<input type="checkbox"/> \$350	<input type="checkbox"/> \$191
Regulation of Combination Products	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$274
Regulation of Dietary Supplements and NHPs	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$274
Regulation of Generic Drugs in the US	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$274
Regulation of US and EU Biologics	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$349
REMS and RMPs	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$274
QUALITY	Member	List	Enterprise
Good Laboratory Practice (GLP)	<input type="checkbox"/> \$365	<input type="checkbox"/> \$500	<input type="checkbox"/> \$274
Good Manufacturing Practice (GMP)	<input type="checkbox"/> \$465	<input type="checkbox"/> \$640	<input type="checkbox"/> \$349
Quality System Regulation (QSR)	<input type="checkbox"/> \$545	<input type="checkbox"/> \$745	<input type="checkbox"/> \$409

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
Medical Devices Postapproval	<input type="checkbox"/> \$1210	<input type="checkbox"/> \$1650	<input type="checkbox"/> \$1089
Regulatory Basics: Complete	<input type="checkbox"/> \$ 500	<input type="checkbox"/> \$ 700	<input type="checkbox"/> \$450
Regulatory Basics: EU	<input type="checkbox"/> \$ 320	<input type="checkbox"/> \$ 440	<input type="checkbox"/> \$288
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
Regulatory Medical Writing: Package #2	<input type="checkbox"/> \$ 970	<input type="checkbox"/> \$1325	<input type="checkbox"/> \$806
Regulatory Medical Writing: Package #3	<input type="checkbox"/> \$ 970	<input type="checkbox"/> \$1325	<input type="checkbox"/> \$806
Regulatory Medical Writing: Package #4	<input type="checkbox"/> \$ 780	<input type="checkbox"/> \$1080	<input type="checkbox"/> \$707

METHOD OF PAYMENT

International Wire Transfer: Fax a completed form and copy of bank wire confirmation to confirm your registration to: 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)