RAPS ONLINE UNIVERSITY
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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 ❑ Home _____ Suite/Apt _____

City/ State/Province _____

Mail Stop _____ Postal Code _____ Country _____

Phone (with area/country code) _____

Email Address _____

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

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

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CUSTOMER NAME

First _____ **MI** _____ **Last** _____

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

REGISTRATION FEES (All fees in US dollars)

BUNDLES	Member	List	Enterprise
Clinical Trial Foundations	❑ $ 575	❑ $790	❑ $518
GxP	❑ $ 865	❑ $1292	❑ $779
Regulatory Basics: Complete	❑ $ 500	❑ $ 700	❑ $450
Regulatory Basics: US and Canada	❑ $ 400	❑ $ 560	❑ $360
Regulatory Medical Writing: Complete	❑ $1240	❑ $1710	❑ $1116
Regulatory Medical Writing: Package #1	❑ $ 1050	❑ $1440	❑ $878

METHOD OF PAYMENT

o **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.

o **Check #**_____

o **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)