RAPS ONLINE UNIVERSITY
Essential knowledge. Well earned.

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 _____

REGISTRATION FEES (All fees in US dollars)

CLINICAL	Member	List	Enterprise
Globalization of Clinical Research Trials and Investigations	❑ $360	❑ $495	❑ $270
Good Clinical Practice (GCP)	❑ $250	❑ $345	❑ $188
Understanding and Managing the US Clinical Trial Process	❑ $455	❑ $630	❑ $341
ESSENTIALS	**Member**	**List**	**Enterprise**
Effective Regulatory Communication	❑ $360	❑ $495	❑ $270
Ethics	❑ $455	❑ $630	❑ $341
FDA Law and Regulation	❑ $455	❑ $630	❑ $341
Intermediate Medical Writing: Pharmaceuticals and Biologics	❑ $360	❑ $495	❑ $270
Intermediate Medical Writing: Investigational Applications	❑ $550	❑ $770	❑ $413
Intermediate Medical Writing: Medical Devices	❑ $250	❑ $345	❑ $188
Introduction to Regulatory Affairs in the EU	❑ $135	❑ $185	❑ $101
Introduction to Regulatory Affairs in the US and Canada	❑ $250	❑ $345	❑ $188
Introductory Medical Writing	❑ $360	❑ $495	❑ $270
Project Management for Regulatory Professionals	❑ $455	❑ $630	❑ $341
Regulatory Due Diligence for Product Development	❑ $360	❑ $495	❑ $270
Role of the Regulatory Professional	❑ $ 25	❑ $ 25	❑ $ 19
Supplier Management	❑ $360	❑ $495	❑ $270
Supply Chain Controls	❑ $360	❑ $495	❑ $270
MEDICAL DEVICES	**Member**	**List**	**Enterprise**
Global Regulatory Strategy for Medical Devices	❑ $455	❑ $630	❑ $341
Medical Devices: Advertising and Promotion in the US	❑ $360	❑ $495	❑ $270
Medical Devices: Canadian Regulations	❑ $360	❑ $495	❑ $270
Medical Devices: China, Japan, Singapore and South Korea Regulatory Overview	❑ $455	❑ $630	❑ $341
Medical Devices: Compliance and Audits	❑ $535	❑ $735	❑ $401
Medical Devices: Corrections, Removals and Directed Recalls	❑ $550	❑ $770	❑ $413
Medical Devices: Definition and Lifecycle	❑ $135	❑ $185	❑ $101
Medical Devices: EU Regulations	❑ $535	❑ $735	❑ $401
Medical Devices: Postmarket Surveillance	❑ $455	❑ $630	❑ $341
Medical Devices: Risk Management	❑ $455	❑ $630	❑ $341
Medical Devices: US Regulations	❑ $535	❑ $735	❑ $401
Regulation of Combination Products	❑ $360	❑ $495	❑ $270
Regulation of IVDs for Key International Markets	❑ $455	❑ $630	❑ $341
Regulation of IVDs in the US	❑ $360	❑ $495	❑ $270

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CUSTOMER NAME

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

PHARMACEUTICALS	Member	List	Enterprise
Chemistry, Manufacturing and Controls	❑ $535	❑ $735	❑ $401
Global Regulatory Strategy for Pharmaceuticals	❑ $455	❑ $630	❑ $341
Pharmaceuticals: Advertising and Promotion in the US	❑ $360	❑ $495	❑ $270
Pharmaceuticals: Canadian Regulations	❑ $550	❑ $770	❑ $413
Pharmaceuticals: Compliance and Audits	❑ $535	❑ $735	❑ $401
Pharmaceuticals: Definition and Lifecycle	❑ $135	❑ $185	❑ $101
Pharmaceuticals: EU Regulations	❑ $550	❑ $770	❑ $413
Pharmaceuticals: US Regulations	❑ $535	❑ $735	❑ $401
Pharmacovigilance	❑ $455	❑ $630	❑ $341
Regulation of Biosimilars	❑ $250	❑ $345	❑ $188
Regulation of Combination Products	❑ $360	❑ $495	❑ $270
Regulation of Dietary Supplements and NHPs	❑ $360	❑ $495	❑ $270
Regulation of Generic Drugs in the US	❑ $360	❑ $495	❑ $270
Regulation of US and EU Biologics	❑ $455	❑ $630	❑ $341
REMS and RMPs	❑ $360	❑ $495	❑ $270
QUALITY	**Member**	**List**	**Enterprise**
Good Laboratory Practice (GLP)	❑ $360	❑ $495	❑ $270
Good Manufacturing Practice (GMP)	❑ $455	❑ $630	❑ $341
Quality System Regulation (QSR)	❑ $535	❑ $735	❑ $401

REGISTRATION FEES (All fees in US dollars)

BUNDLES	Member	List	Enterprise
Clinical Trial Foundations	❑ $ 1000	❑ $1370	❑ $900
GxP	❑ $ 855	❑ $1175	❑ $770
Medical Devices Postapproval	❑ $1210	❑ $1650	❑ $1089
Regulatory Basics: Complete	❑ $ 500	❑ $ 700	❑ $450
Regulatory Basics: EU	❑ $ 320	❑ $ 440	❑ $288
Regulatory Basics: US and Canada	❑ $ 400	❑ $ 560	❑ $360
Regulatory Medical Writing: Complete	❑ $1164	❑ $1632	❑ $1048
Regulatory Medical Writing: Package #1	❑ $ 1050	❑ $1440	❑ $878
Regulatory Medical Writing: Package #2	❑ $ 970	❑ $1325	❑ $806
Regulatory Medical Writing: Package #3	❑ $ 970	❑ $1325	❑ $806
Regulatory Medical Writing: Package #4	❑ $ 780	❑ $1070	❑ $652

METHOD OF PAYMENT

o **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.

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: RAPS.org/onlineu (credit card only)
MAIL: RAPS c/o SunTrust Lockbox Dept, PO Box 79546, Baltimore, MD , 21279-0546
FAX: +1 301 841 7956 (credit card or wire)