DIA IASMED

Protocol Developemt

4 September, 2026

7:30 AM – 11:30 AM | Indonesia, Thailand, Vietnam, Cambodia, Laos
8:30 AM – 12:30 PM | Singapore, Malaysia, Philippines, Hong Kong, Taiwan
Zoom Webinar

Designing High-Quality Clinical Trial Protocols

Designed for clinical research professionals involved in protocol development, study planning, and clinical trial execution, this interactive online training provides the knowledge and practical skills required to design scientifically sound, ethically appropriate, and operationally feasible clinical trial protocols.

Conducted in collaboration with IASMED, the program combines international best practices, practical guidance, and real-world examples. Through interactive discussions and expert insights, participants will learn how to translate research concepts into well-designed clinical protocols that support regulatory compliance, efficient study execution, and successful clinical research outcomes.

Key Learning Topics

- Overview of Clinical Research and Drug Development
- Ethics in Clinical Research
- Protocol Development
- Planning and Conducting Clinical Research

Who should attend?

- Clinical Research Associates (CRAs)
-Clinical Research Managers
- Clinical Trial Project Managers
- Clinical Scientists
- Medical Affairs Professionals
- Sponsor Representatives
- CRO Professionals
- Regulatory Affairs Professionals
- Clinical Operations Professionals
- Study Coordinators
- Investigators & Sub-Investigators
- Research Nurses
- Academic Researchers

Live Online Training with Recording Access

Registered participants will receive access to the session recordings, allowing them to review the content or catch up on any sessions they are unable to attend live.

To register, please visit the link below or scan the QR code.

REGISTER NOW

FACULTY INFORMATION

Lynn King, MHA
Consultant,
Cornerstone Clinical Research Consulting,
United States

- 25+ Years of Experience in Clinical Research & Drug Development
- Extensive Experience Across Pharmaceutical Companies, CROs & Research Sites
- Expert in GCP, Clinical Trial Operations & Regulatory Compliance
- Certified Trainer & Professional Coach

DIA MEMBERSHIP

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DIA volunteers, members, and staff provide a comprehensive catalogue of conferences, workshops, training courses, scientific publications, and educational materials, throughout the year, all around the world.

DIAglobal.org

DIA Korea
Korea@DIAglobal.org

AGENDA | Protocol Development - 4 September, 2026

Session Agenda	DAY 1 (4 September)
Welcome & Objectives	Welcome & Objectives • Learning objectives review • 3-5 question poll
Module 1: **Drug Development Overview**	• Drug Development: Industry trends review • Clinical Development Plans • Regulatory oversight and influence in protocol development
Module 2: **Patient and Provider Perspectives**	• Patient perspective in clinical protocol development • Evaluating context for clinical practice and provider input • Assessing marketplace and standard of care influence on protocol strategy
Module 3: **Protocol Development Process**	• Review typical process for protocol strategy and development • Overview of roles involved in protocol writing, review and finalization • Comparison of Sponsor and Investigator-initiated trials
Module 4: **Protocol Design Overview**	• Sections of a clinical trial protocol • Key factors to a successful protocol design • Considerations for operationalizing a protocol design • Protocol amendment discussion
Module 5: Protocol Case Study	• Review prior protocol designs and discuss alternatives • Discuss challenges in protocol design for sites and patients
Q&A	• Live Q&A, participant-submitted questions addressed

REGISTRATION FORM : Register online or forward to DIA Korea Korea@DIAglbal.org

DIA Webinar

Protocol Development
4 September 2026 | Zoom Webinar

REGISTRATION

Register online at the link HERE or complete this registration form and email to our Korea Office

DIA will send participants a confirmation letter within 10 business days after receipt of their registration.

Registration Fee: USD 100

The registration fee for this course is the same regardless of DIA membership status.Registration fee includes full admission to virtual course, electronic access to training course materials. Please note that the full amount must be received by DIA by commencement of the course to get the electronic access to the material. If you wish to register for both the Protocol Development Training and the Advanced GCP for Clinical Trial Professionals course, please select "Both Trainings" from the table below, complete this registration form, and email it to our DIA Office Korea@DIAglobal.org.

	REGISTRATION FEE (USD)
2-3 Sep | Advanced GCP for Clinical Trail Professionals	☐ 150
4 Sep | Protocol Development	☐ 100
2-4 Sep | Both Trainings	☐ 250

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☐ I DO NOT want to be a DIA member

To view the benefits of the membership, please click the link.

Please check the applicable category:

☐ Academia ☐ Government ☐ Industry

Last Name

First Name

Degrees ☐ Dr. ☐ Mr. ☐ Ms.

Job Title

Company

Address (As required for postal delivery to your location)

City State Zip/Postal Country

email **Required for confirmation**

Phone Number **Required**

CONTACT INFORMATION

DIA
email:
Korea@DIAglobal.org
Eunah.Cha@DIAglobal.org
www.diaglobal.org

DIA Terms and Conditions

CANCELLATION POLICY: On or before 2 August, 2026 Administrative fee that will be withheld from refund amount: the administrative fee that will be withheld from refund amount is 25 % of the delegate fee

Cancellations must be in writing and be received by the cancellation date above. Registrants who do not cancel by that date and do not attend will be responsible for the full registration fee paid.

You may transfer your registration to a colleague at any time but **membership is not transferable**. Please notify DIA of any such substitutions as soon as possible. Substitute registrants will be responsible for nonmember fee, if applicable.

EVENT STREAM AND RECORDING

If you attend a DIA event, we make video and audio recordings of events (both face to face and online) that may include your participation in the event, including your image, questions and comments. To view our full photography and video recording policy, click **here**. (https://www.diaglobal.org/general/photography-policy)

PRIVACY STATEMENT

The personal information you request will be used for the purpose of sending conference information from DIA. In addition, in the web conference, we will use the information with the name of the company or organization and the name of everyone who participates, and it will be used for networking with participants, related parties, exhibiting companies for the period and about two weeks after the event. By submitting this application form, it is interpreted that you have consented to the above handling of personal information, but if you do not agree, please contact DIA.

By signing below I confirm that I agree with DIA's Terms and Conditions of booking. These are available from the office or online by clicking **here**. (https://www.diaglobal.org/general/photography-policy)

Signature Date

PAYMENT OPTIONS

Register online at **www.DIAglobal.org** or check payment method.

☐ **BANK TRANSFER:**

You will recieve an invoice with bank information detail by email after registration completion.
All local and overseas charges incurred for the bank transfer must be borne by payer.

☐ **CREDIT CARD (VISA OR MASTERCARD OR AMEX ONLY)**

☐ **VISA** ☐ **MC** ☐ **AMEX** Exp. (mm/yy) _____

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