DIA

Global Labeling Conference

WHO

Professionals from biopharmaceutical and device companies, regulatory authorities, CROs, and consulting agencies involved in:

- Labeling
- Clinical safety/pharmacovigilance
- Pharmacoepidemiology
- Regulatory affairs/drug review and approval process
- Medical affairs and communications
- Medical writing
- Clinical research and development
- Product research and development alliances
- Quality control/quality assurance

WHAT

DIA's Global Labeling Conference

WHEN

April 24-25

WHERE

North Bethesda, MD

WHY

- Discuss new labeling-related developments and regulations in diverse global regions such as Canada, EU, Japan, Asia, Latin America, Middle East, Africa, and US
- Describe the impact of proposed changes to regional and global labeling requirements and implications for labeling practice and processes
- Analyze the impact of current and proposed global and region-specific labeling policies for combination products, labeling development, and product lifecycle practices
- Assess the advantages and limitations of various labeling processes and document management systems commonly used in companies
- Describe global labeling compliance expectations and best practices

SECURE YOUR SEAT