The Sunshine Act: Reporting for Clinical Trials

Description: The Sunshine Act or Open Payments Program requires manufacturers of drugs, medical devices, and biologics that participate in U.S. federal health care programs to report certain payments and items of value given to physicians and teaching hospitals. This Act was part of a healthcare reform bill adopted in March 2010. It came about due to requests for increased transparency about the financial relationships between physicians and industry. The Centers for Medicare and Medicaid (CMS) issued the final rules in 2013 which implemented the Sunshine Act.

Objectives of the Presentation:
- Purpose of the Sunshine Act
- Who is required to report under the Sunshine Act?
- What is reported?
- Exclusions
- Tracking
- Penalties
- Useful links

Why Should you Attend:

The Sunshine Act exposes the physicians and sponsors to new reporting requirements which are unprecedented in their scope and detail noncompliance to which would lead to hefty fines. This webinar will summarize the new requirements and provide practical solutions to most common situations that are prevalent and necessary between sponsors and investigators. Templates for reporting format, acceptable accounting practices, exemptions to reporting requirements, and timelines for compliance and enforcement will be discussed.

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