



Presents

SCABBinar:

Biological Product Deviations: Issues on the Front Lines

Tuesday, January 13, 2009

Presented by:

Glenn Ramsey, MD

Description: The FDA began requiring biological product deviation (BPD) reporting in 2001, and provided additional guidance in 2006. This presentation will focus on the most common and the most serious BPDs, especially for transfusion services, and what trends have developed over recent years. BPDs provide valuable lessons for compliance and quality improvement and in some cases represent potentially serious risks for transfusion recipients.

Objectives:

- 1) Review five-year national trends in biological product deviations.
- 2) Discuss the most common compliance requirements involved.
- 3) Consider the implications for transfusion safety.

Level of difficulty: Intermediate

Audience: Technologists, supervisors, laboratory directors and physicians

PACE Contact Hours – 1

Tuesday, January 13, 2009

9:00 am PDT & AZ, 10:00 am MDT, 11:00 am CDT, 12:00 pm EDT

Registration Fee:

Individual Member: \$25.00

Individual Non-Member: \$80.00

Institutional Member: \$50.00

Institutional Non Member: \$100.00

The SCABBinar is facilitated via live conference call. All handouts, evaluations and attendance rosters will be provided for download prior to the SCABBinar.

To register, visit www.scabb.org > Education Events