The CRC's Guide to Coordinating Clinical Research

Karen E. Woodin, Ph.D.

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The CRC's Guide to Coordinating Clinical Research is a comprehensive training resource for investigative site staff. This invaluable guide offers CRCs the information they need to successfully coordinate a clinical trial from study startup to closeout and beyond.

Topics covered include:
*Developing standard operating procedures (SOPs)
*Recruiting and retaining study subjects
*Understanding the informed consent process
*Working with protocols and case report forms
*Recognizing adverse events
*Preparing for audits

The CRC's Guide to Coordinating Clinical Research is recommended for:
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