CenterWatch Announces — Clinical Trial Billing Documentation Best Practices: Optimize Reimbursement Webinar, April 24, 2019

Billing and reimbursement are key to the bottom line success of trials. Optimize the reimbursement with best practices.

FALLS CHURCH, Va. (PRWEB) April 12, 2019 -- Clinical Trial Billing Documentation Best Practices: Optimize Reimbursement
**A CenterWatch Webinar**
Wednesday, April 24, 2019, 11:00 a.m. - 12:30 p.m. EDT

The devil is in the details. And reimbursements require lots of details. Don’t lose money by not complying with Medicare regulations, FDA guidance or industry best practices.

Nancy Reynolds Howard, RN, BSN, CCRC — consultant at NRH Compliance Partners — will discuss the nuances and best practices for optimizing reimbursement through documentation of medical necessity, documentation of treatment provided to the subjects, coding of services provided and conducting an annual review of your key research documents.

During this webinar attendees will learn how to collect the maximum reimbursement due. Nancy will cover the:

• Relationship between documentation of medical necessity and billing compliance
• Enhancement of billing compliance using medical necessity language and document consistency reviews
• Connection between the language used in the informed consent form and billing compliance
• Nuances of specific codes and modifiers for claims under Medicare’s clinical trial policy and investigational medical device regulation
• Importance of complying with Medicare diagnostic and procedural codes: diagnostic code Z00.6 — Examination of participant in clinical trial; procedure code modifier Q0; and procedure code modifier Q1 — “Routine Costs”

Secure the monies for services provided. Optimize reimbursements through best practices in billing, documentation and compliance. Visit our website for more information about the speaker, pricing, and more.

Interested in registering multiple sites?
Call (888) 838-5578 in the U.S. or +1 (703) 538-7600 globally to learn about our special multisite discount.

Webinar Details:
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PRWeb ebooks - Another online visibility tool from PRWeb
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By phone: 617-948-5100 or 866-219-3440

About CenterWatch:
Founded in 1994, CenterWatch is a trusted source and global destination for clinical trials information for both professionals and patients. CenterWatch provides proprietary data and information analysis on clinical trials through a variety of newsletters, books, databases, and information services used by pharmaceutical and biotechnology companies, CROs, SMOs, and investigative sites involved in the management and conduct of clinical trials. As a pioneer in publishing clinical trials information, CenterWatch was the first Internet site to publish detailed information about active clinical trials that could be accessed by patients and their advocates.
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You can read the online version of this press release here.