

IVRS in Clinical Trials (IVRS-01)

29 April 2014, Berlin



IVRS & Logistics Training

Explore the Capabilities of IVR(S)-Systems and what Advantages can be gained by using this Technology!



Instructor

Nimer Yusef, PMP





Content

- ☀ IVRS-Overview
- ☀ IVRS Functions
- ☀ Benefits / Potential
- ☀ Processes

Limited to 12 Attendees






Training Objectives

This Training will provide you with a practical Introduction into the World of IVR(S)-Systems, sometimes also referred as IRT-Systems. By attending the Course, you will gain of the Procedures and Techniques needed to reach the optimal Functionality of these Systems.

-  Comparison of IVRS and non-IVRS Studies
-  Functions and Functionality of IVR(S)-Systems
-  Randomization using IVRS
-  Compliance with regulatory Requirements via Implementation of IVRS

Who should attend?


The Training is designed to benefit Clinical Trial Professionals, to Specialists and Managers within Pharma Companies and CROs. Target audience is:

-  Clinical Project Manager / Monitor / CRA
-  Clinical Logistics - / Clinical Trial Supply Manager
-  QA-Manager / Auditor
-  Consultant
-  IVRS-Specialist

Newsletter

If you are interested in being informed of further trainings, please subscribe to my Newsletter. (E-Mail: Newsletter@trial-brain.com).

Instructor

 Nimer Yusef, PMP

is a Mathematician and certified Project Manager (PMP) with a strong background in Information Technology. His first experiences within the clinical research industry he gathered with the CRO ICRC Weyer more than 10 years ago. Subsequently, he worked for Perceptive Informatics, where he was the responsible Project Manager for IVRS-System Development with a focus in the area of Inventory Management. As Technical Logistics Leader at Parexel Clinical Logistics Services he is responsible for training and consulting in IVRS-technology concepts since 2008. To close the competence gap between IVRS providers and end users in clinical study teams he founded the Company Trial Brain in 2010 to offer Seminars, Trainings and Workshops for IVRS and Clinical Logistics.

Date

Tue 29-Apr-2014, 09:00 AM to 05:00 PM

Location

Einstein Palais Friedrichstraße, Berlin
near the Station Friedrichstraße

Costs

900 Euro + VAT per attendee (includes documentation and catering).

Register 1 month in advance for a 10% rebate!

History of IVRS

- ☀ Blocked Randomization
- ☀ Site Stratification
- ☀ Development from early Approachs through to contemporary IVR(S)-Systems
- ☀ IVRS Demo

IVRS Functions

- ☀ Typical Scope of Functionality
- ☀ Users and Roles within the System
- ☀ Limitations and special Features
- ☀ Processes in correlation with IVRS
- ☀ Benefits and Drawbacks of using IVR(S)-Systems

Integration

- ☀ Differences between IVRS-Providers
- ☀ Selection of an IVRS-Provider
- ☀ Integration into internal / external Systems
- ☀ User Training
- ☀ Preparation & Execution of User Acceptance Testing (UAT)

Randomization

- ☀ Randomization Fundamentals
- ☀ Practical Use of Stratification and Randomization Blocks
- ☀ List-based Randomization
- ☀ Dynamic Randomization
- ☀ Pratical Exercise in Randomization

Inventory Management

- ☀ Inventory Management Fundamentals
- ☀ Separation of Patient & Medication-randomization
- ☀ Blinding of Depots and Sites
- ☀ Fundamentals of Threshold & Prediction based supply Methods
- ☀ Processes in Conjunction with Inventory Management

Benefits of using IVRS

- ☀ Process Implementation & Compliance Support
- ☀ Track & Trace, Reporting

Registration

Details

Fax: +49 / 30 / 81 45 84 99

E-Mail: events@trial-brain.com

Online: at trial-brain.com

Registration

Registration for the Seminar "IVRS in Clinical Trials" (IVRS-01) 29-Apr-2014 in Berlin, Germany.

Name

Position

Department

Company

Street

Postal Code / City

Country

Phone

E-Mail

Notes

Date & Signature

Date & Time

Tue 29-Apr-2014, 09:00 AM to 05:00 PM

Location

Einstein Palais

Friedrichstraße 171

10117 Berlin

Accommodation

There is no special offering for participants of this seminar. The hotels [The Westin Grand](#) & [Hilton](#) can be recommended.

Costs

900 Euro + VAT per attendee (Fee includes Documentation and Catering).

Register 1 month in advance for a 10% rebate!

Cancellations by Registrants will be associated with the following costs:

- until 1 Month, 100 Euro + VAT

- until 2 Weeks, 400 Euro + VAT

A Substitution of a Registrant is possible anytime free of Charge.

Contact

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