

SEMINAR

Standards In Electronic Data Capture

Tuesday 1st November 2005

DTI Conference Centre
1 Victoria Street
London SW1H 0ET

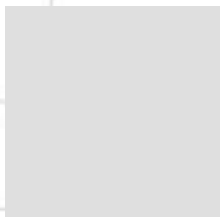


SUMMARY OF CONTENT

Standards are being implemented at the back end of EDC (CDISC, HL7) but not currently at the front end, the user interface. This is an area that receives less attention than it should. Drawing on the extensive experience of the Panel, the four presentations will discuss key aspects of any EDC study and will highlight areas where there are challenges and their potential solutions. During the Panel question and answer session, you will have the ability to tap into this experience to assist with your approach to EDC and its standardisation.

SPEAKERS AND CHAIRS

Dr Emma Banks	Head of Data Services, Datatrial Ltd
Mr Birant Osman	Group Manager, Clinical Data Manager, Statwood Ltd
Dr Bill Byrom	Director of Product Strategy, ClinPhone Group Ltd
Mr Andrew MacGarvey	Vice President (Europe), Datatrial Ltd
Mr Rob Nichols	Director of Data Operations & Business Systems, GDRU Quintiles Ltd
Mr Anil Vaidya	Biotechnology & Pharmaceuticals Sector Specialist, UKTI



PROGRAMME

1.30 Registration

2.00 Introduction

Anil Vaidya & Andrew MacGarvey

2.20 The Impact of EDC on Current Monitoring and Site Roles

Birant Osman

Abstract: This presentation aims to explore the impact of EDC on smaller and medium sized CROs and monitoring practices; the changes in the role of monitors from the sites perspective and the changes in roles for site personnel themselves. Any associated problems with these issues, and their possible resolutions, will also be discussed.

2.50 New Role Models for the EDC Study Team – Training, Workflow & Service Provision

Emma Banks

Abstract: With the increased uptake of EDC in the clinical trials arena, the roles of key team members will have to evolve to embrace the new technologies and workflows. The presentation aims to define the potential evolution of roles within the EDC study team and how workflows can be managed and standardised whilst maintaining a high level of service provision.

3.20 TEA

3.50 How to Support the 200 or so Different EDC Systems Currently Available

Rob Nichols

Abstract: The speaker will present a case study outlining how the GDRU Quintiles Phase I clinical unit, working each year on over 60 different studies, plans to manage the expected increase in EDC studies. Key has been the development of an in-house solution while maintaining a flexible approach to working with clients own EDC solutions.

4.20 An Integrated Approach to Successful eClinical Trials

Bill Byrom

Abstract: This presentation will explore the key benefits of combined data sources and integrated solutions. Case studies and practical demonstrations will be presented to illustrate the benefits of integrating Interactive Voice Response (IVR), Electronic Data Capture (EDC) and Clinical Trial Management Systems (CTMS) – providing Sponsors and end-users with the tools necessary for enhanced trial management processes.

4.50 Panel Discussion

5.15 Cold Fork Buffet with Wine

COST:

The cost of the Seminar (including buffet) is £60 + VAT (CCRA members) or £90 + VAT (non-members).

TO BOOK:

Please email your booking request (giving number of places required and details of invoice address) to mail@ccra.org.uk. Telephone bookings will also be received on 0116 271 9727.



For more detailed information on how membership of the CCRA can specifically benefit your organisation, please contact us through any of the channels below:

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