

Data Transparency Autumn Event 2024



Save the Date!

The Data Transparency Autumn Event will take place from 17–19 September 2024. This virtual event provides a platform for attendees and presenters to foster meaningful interactions and collaborate on ideas. Presentations will be delivered across the three days in bitesize chunks from **1 5:00-17:30 (BST) / 10:00-12:30 (EDT)**.

Call for speakers is now open. Don't miss out on the opportunity to contribute your expertise, innovations, and research by submitting your **150-word abstract** before **14 June**.

Need guidance? Explore the curated topics below from PHUSE's Data Transparency Leads to spark inspiration for your standout abstract.

- Data Anonymisation Techniques and Experiences
- Risk Quantification
- Data Sharing Initiatives and Processes
- Registries and Results Reporting
- Synthetic Data
- Data Transparency Regulatory Submissions
- Plain Language Summaries (PLSs) and Plain Language Summaries of Publications (PLSPs)
- EU CT

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Save the date for three days of engaging discussions and insightful presentations! Visit the [PHUSE archive](#) for all previous event recordings and presentations.

Data Transparency Autumn Event Sponsors

Virtual Event Sponsors



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Sponsor Flyers

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Simplify clinical trial transparency with our unparalleled expertise

Recent Global Data Transparency initiatives have pushed the clinical trial disclosure scope ever wider to include not only registration and results postings, but also sharing of de-identified clinical trial documents, and the delivery of clinical results in a plain language format.

Regulatory agencies, industry trade organizations, and a growing base of global stakeholders continue to work to increase the breadth of trial information being made available in response to the rising demands from both public and private spheres alike.

CTG Registration

Extract Data

Details

Send to Author

13 Nov 2022

26 Nov 2022

6 Dec 2022

14 Dec 2022

22 Dec 2022

1 Jan 2023

Preparation

Author

QC

Review

Approve

Expert services to help you navigate through global transparency with confidence

Clinical Trial Disclosure

Comprehensive and expert support for accurate and timely compliance of global transparency obligations.

Medical Writing & Plain Language Summary

Development of full suite of clinical trial documentation from early phases of development to trial completion.

Transparency

Fulfillment of de-identification requirements through machine learning and pattern searching.

Advisory Services

Helping sponsors navigate the evolving transparency world through advisory services based on experience and best practices.

Solutions Overview

3 Ways to Safely Accelerate Your Clinical Document Submissions

Enabled by powerful privacy technology

PRIVACY ANALYTICS

an IQVIA company

What you're up against

As a clinical trial sponsor, you're facing pressure from multiple sides when it comes to handling regulatory requirements for document anonymization and redaction.

Transparency

Under EMA Policy 0070 and Health Canada PRCI, safety preserving statistical anonymization of personal data is preferred for transparency.

Commercial protection

You're under increasing pressure to protect commercially confidential information (CCI) as well as participant privacy.

Timelines

Timelines remain tight, especially with retrospective requests, and are highly restrictive under the EU Clinical Trials Regulation.

Cost-efficiency

Manual approaches can be resource intensive, and you often face unexpected spikes in demand for publication.

Meet tight timelines, ensure a competitive edge

To help you ensure a competitive edge, Privacy Analytics offers three paths to protecting CCI and personal data while meeting deadlines for EU CTR and other initiatives such as Health Canada PRCI and EMA Policy 0070: **Redaction-as-a-Service, Anonymization-as-a-Service, and licensed software.** Each solution is powered by our 4th-gen document redaction and anonymization software platform.

With Privacy Analytics by your side, you'll be better able to safely accelerate submissions for the broader range of clinical document types you now need to address throughout the clinical trial life cycle.

Reduce time to submit clinical trial documents from weeks to days

Our latest software release unites our proven natural language processing technology with automated redaction, providing you with a seamless solution to meet all of your evolving data privacy goals.

Sponsorship

Hosting the Data Transparency Events digitally means that no matter where you are in the world you can participate. It provides the industry with a broader opportunity to share knowledge on a global scale, connecting through the virtual event platform. The sponsor options offer a range of benefits with ample company exposure. See the prospectus for more detail.

phuse.digital

Data Transparency Events

Winter Event: 6–8 February 2024
Autumn Event: 17–19 September 2024

Prospectus

phuse.global

Data Transparency Working Group Leads



Stephen Bamford
*Janssen Research &
Development*



Jean-Marc Ferran
Qualiance



Devaki Thavarajah
Independent



Muhammad Oneeb
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