DOMINIKA
DATA PROTECTION COORDINATOR LIFE SCIENCES

SPECIALISMS

- Clinical Trials
- Healthcare
- Life Sciences
- NHS Data Security and Protection Toolkit (DSPT)
- Hospitality

LANGUAGES

- Polish (native)
- Czech/Slovak
- English

PROFILE

Dominika has a strong background in clinical trials management and a BSc (Hons) in Health Sciences. With hands-on experience across the full clinical trial lifecycle, she brings a practical understanding of regulatory requirements, ethical considerations, and risk management in complex research environments.

Dominika's experience designing and managing multi-jurisdictional and paediatric trials gives her a strong foundation in handling sensitive personal data and navigating sector-specific regulations. She supports Life Sciences organisations by applying her analytical and collaborative skills to deliver compliant, well-structured data protection processes.

QUALIFICATIONS

- Essentials of Clinical Research Monitoring
- Advanced Project Management in Clinical Trials
- Data Management in Clinical Research

SAMPLE EXPERIENCE

As a Clinical Trial Manager within university research centres, Dominika led operational and quality aspects of clinical studies, ensuring compliance with ICH-GCP standards and regulatory requirements. She supported the development of Informed Consent Forms (ICFs), protocol-specific documentation, and ethics submissions, whilst maintaining trial master files and coordinating cross-functional trial teams.

Dominika also conducted comprehensive risk assessments at the outset of each study, developing mitigation plans and implementing ongoing monitoring to maintain trial integrity. She has extensive experience managing paediatric and complex therapeutic area trials, where ethical considerations, patient safety, and regulatory compliance were central to study design and delivery.