Who you'll meet at the **Clinical Trial Regulation and Clinical Data Transparency** Conferences

6-8 December 2016 | London, United Kingdom

Account Manager
Ass. Director, Global Clinical Trial Submission Unit GCT-SU and EU CT External P
Associate Director CTA
Associate Director GRA CMC
Associate Director, Global Clinical Study Disclosure
Chief Policy Adviser
Chief Strategy Officer
Cinical Trial Assistant
Clinical Assessor
Clinical Compliance Manager
CLINICAL OPERATION SPECIALIST
Clinical Quality Assurance Associate
Clinical Study Manager
Clinical Trial Administrator
Clinical Trial Assistant
Clinical Trial Disclosure, Clinical Development
Clinical Trial Regulatory Affairs Manager
Consultant
Department Head
Dir. Gloabl R&D and GVP QA
Director
Director Clinical Operations
Director Global Regulatory & Scientific Policy
Director of Clinical Trials & Regulatory Services
Director Regulatory Affairs , EFPIA, Belgium
Director Regulatory Policy and Intelligence
Director, Clinical Trials Regulatory Group
Director, Clnical Trial Information Disclosure
Director, Data Transparency Port.Off.
Director, Global Clinical Registry
Director, Office of Medical Transparency
Director, Policy & Intelligence
Director, Policy, Chief Medical Office
Divisional Director
EU Regulatory Policy Mgr & Clinical Ops Quality Mgr
Event Manager
Executive Director, Global Regulatory Policy
Exhibition Manager
Expert Inspector, GCP
Global DSUR Manager
Global Head Central Solutions and Services Medicine (CSSM) in Global Biostatisti
Global Head of Medical Writing and Disclosure
Global Scientific Information Manager
Head Clinical Research
Head of Clinical and Non-Clinical Compliance
Head of Compliance and Inspections Department
Head of International Regulatory and Intergroup Office
Head of the Clinical Disclosure Office
Head Regulatory & Matrix Services
Manager
Manager, Clinical Trial Transparency
Manager, Regulatory Affairs
Managing Editor
Medical Writer
Nonclinical Assessor, Clinical Trial Unit
Policy Director
Principal
Principal Advisor - Global Regulatory Affairs Europe
Principal Medical Writer
Programme Director Technology Appraisals, PASLU and HST
Project Manager
Public Disclosure Lead
QA Manager
Quality Assurance Manager
RA Liaison Specialist
Reader in Medical Statistics
Regulatory Affairs Associate Director
Regulatory affairs Director - EU Policy
Regulatory Affairs Manager
Regulatory Affairs Manager
Regulatory Affairs Manager & Safety Advisor
Regulatory Affairs Officer
Regulatory Affairs Project Manager
Regulatory Affairs, Team Leader- Speciality Care
Regulatory Consultant
Regulatory Manager
Regulatory Professional
Regulatory Scientist
Scientific Collaborator
Senior Clinical Operations Manager
Senior Director, Global Regulatory Affairs & Policy, Europe
Senior Director, Worldwide Regulatory Strategy
Senior GCP Advisor
Senior GCP Advisor
Senior Manager of Regulatory Affairs
Senior Manager Regulatory Affairs
Senior Partner
Senior Regulatory Affairs Manager
Senior Reporter, Scrip Regulatory Affairs
Senior Trial Manager
Specialist CTA
Sr. Expert Clininal Trial Disclosure and R&D Processes
Vice President
Vice President, Global Regulatory Affairs, Oncology
Vice President, International Regulatory Affairs

Sign up today to ensure your space!

Contact Anna.Silva@DIAglobal.org for more information.