AGENDA

BRAND NEW Medical Devices Roundtable Forum 2021

3rd February 2021

All times are in CET

Attend the only event dedicated to bringing delegates together to share best practice and develop solutions by taking part in up to 3 interactive roundtable discussions. As a delegate you'll get a choice of 10 roundtables to choose from.

The Medical Devices Roundtable Forum offers a unique opportunity to come together with your peers in a virtual environment. Hosted by industry experts and each focused on a single issue, roundtables are an exciting, interactive way to build your personal network and learn from the experience and expertise of others.

Places are limited on each roundtable so register your interest today.

3rd February 2021	
08:45	*Registration*
08:55	**Opening address from GlobalData**
09:00	**OPENING KEYNOTE: Introduction and overview to the European medical device industry in 2021** **John Shillingford,** Director of Clinical Research, **Afon Technology**
09:30	**Direct-to-Patient Clinical Trials and the unique considerations for Medical Devices** *Speaker TBC*
10:00	**PANEL DISCUSSION: The impact of the new MDR regulation on the European medical device industry** • Preparing for MDR: what do you need to take into account? • Best practice for meeting the MDR requirement • Navigating unknowns and clarifying new requirements of MDR • Writing and structuring MDR documents to ensure critical content is covered • What are the exact requirements on data to be presented for certification process to notified bodies? *Panellists TBC*

11:00	**Roundtable round 1**
12:00	**Roundtable round 2**
13:00	**Roundtable round 3**
14:00	**Meeting Allocation**
14:30	**CLOSING KEYNOTE: Projections for Medical Device Clinical Trials in 2021** **Robin Sutherland,** Head of Human Resources & Clinical Operations, **Renovia**

Roundtable Titles:

Each table runs 3 times on the same topic. Tables are exclusive, other vendors cannot attend.

Attendees can select up to 3 roundtables.

1. Choosing the appropriate PMCF activity, study, registry or survey? - **SMART-TRIAL (Medei)**
2. The usage of real-world evidence for medical device trials - **IQVIA MedTech**
3. Is the UK still a destination of choice for clinical evaluation? - **NIHR**
4. Clinical Evaluation and Post Market Clinical Follow-up under MDR – basis and practical cases - **Qmed Consulting**
5. Regulatory Evolution in Medical Device Development – **Fiona Maini,** Principal, Global Compliance and Strategy, **Medidata**
6. Roundtable hosted by **Evnia**
7. Roundtable hosted by **Lumis**

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