26 January 2024

Come meet the DKMA and let's talk about developments within regulatory affairs in Europe!

Invitation to Medicoindustrien' s Expert Groups for: Regulatory affairs, quality assurance, clinical investigations and consultants. Hybrid meeting, however, only app. 45 seats in the room, the rest on Teams. See Teams link & register here for the event:

Monday 26 February 2024, 14.00 – 16.00 at

Medicoindustrien / Medtech Denmark, Bøge Allé 5, 2970 Hørsholm

Agenda

14.00 – 14.10	**Welcome & introduction** by Lene Laursen, Medicoindustrien
14.10 – 15.10	**DKMA: Setting the European scene** Latest proposal from the EU-Commission re MDR&IVDR MDCG & CAMD Expert Panels, HERA-matters & EMA involvement Q&A By Katrine Bering Klausen, Senior Advisor, public and international affairs, & Stine Jønsson, Head of Unit, External Collaboration & Coordination, Danish Medicines Agency, Medical devices
15.10– 15.30	**Coffee break**
15.30 – 16.00	Continued discussion
16.00 – 16.15	**Wrap up, Q&A and closing of the meeting**