Canada

Medical Device Regulation in the European Union - Industry Essentials

On October 7, 2020 the Trade Commissioner Service with Global Affairs Canada hosts an exclusive webinar with the **EU Competent Authority for Medical Devices/Danish Medicines Agency** and the consultancy **AlfaNordic.**

The webinar addresses regulatory and practical aspects to consider for medico companies looking to expand their business to Europe. This webinar is targeted and relevant for Canadian executives with plans of activities in Europe within 1-2 years.

Programme:

17:30 PM*	**Welcome and Introduction to European based Trade Commissioners** Jakob Schmidt, Embassy of Canada to Denmark, Trade Commissioner, Life Science Aurora Polo, Consulate General in Barcelona, Life Science, Sector Team for Europe
17:35—17:55	**Medical Device Regulation in the EU before and after May 26th 2021** *Legislation and regulatory framework on medicinal devices in Europe* **Thomas Wejs Møller**, Head of the EU network Competent Authorities for Medical Devices, CAMD and Section Manager - Medical Devices, Danish Medicines Agency
18:00 – 18:20	**How hard can it be? Practical experiences with implementation** *Translating regulatory requirements into every-day business: Typical obstacles and challenges.* **Ole Markersen,** Director - Medical Devices, AlfaNordic
18:20 – 18:40	**Questions and answers** **Thomas Wejs Møller,** The Danish Medicines Agency and **Ole Markersen,** AlfaNordic
18:40 – 18:45	**Vote of thanks and closing remarks** Jakob Schmidt, Embassy of Canada to Denmark, Trade Commissioner, Life Science

*All time points are specified according to local time, Copenhagen, Denmark: Central European Summer Time, UTC+2.

For call-in details and registration, please get in touch with Trade Commissioner Jakob Schmidt: jakob.schmidt@international.gc.ca. RSVP no later than September 28, 2020.

Get in touch for more information and registration

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