U.S. Commercial Service Webinar

Canada: How to Prepare for the Medical Device Single Audit Program (MDSAP)

What is MDSAP?

The Medical Device Single Audit Program (MDSAP) is a global initiative that allows the conduct of a single regulatory audit of a medical device manufacturer's quality management system that satisfies the requirements of multiple regulatory jurisdictions. The MDSAP is a way that medical device manufacturers can be audited once for compliance with the standard and regulatory requirements of up to five difference medical device markets: Australia, Brazil, Canada, Japan and the United States.

Why is MDSAP Important to my company?

All medical devices in Canada must be deemed safe and effective, including software accompanying any medical device. Beginning January 1, 2019, U.S. companies marketing Class II, III or IV devices in Canada will need a valid MDSAP certificate to get, maintain or amend a medical device. In this webinar, *Frédéric Hamelin, Manager of the Quality Systems Section of the Medical Devices Bureau at Health Canada*, will provide an overview of the MDSAP requirements and how to prepare for the transition. This webinar will discuss:

- The status of the MDSAP Transition
- Are companies ready?
- The process to become MDSAP certified

About the U.S. Commercial Service:

Located across the United States and in U.S. Embassies and Consulates in over 70 countries, our global network of trade professionals helps U.S. businesses succeed in markets around the world. Whether you're looking to make your first export sale or expand to additional international markets, we offer the trade counseling, market intelligence, business matchmaking, and commercial diplomacy you need to connect with lucrative business opportunities.

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Event dates

Thursday, June 28, 2018
2:00-3:00 pm EST

Location

Webinar via your computer & telephone

Cost

U.S. $25.00

More information

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