

Medical Device Roundtable: Post-Market Surveillance Obligations

November 18, 2021

From medical device manufacturers and importers to hospitals and health professionals, in this webinar, the panel of regulatory and legal experts covered the following:

- 2021 updates to the Medical Devices Regulations on post-market surveillance obligations of medical device license holders;
- Reporting obligations to Health Canada on adverse incidents involving medical devices sold and imported into Canada; and
- Use of post-market surveillance regulatory disclosure in medical device litigation and cross-border issues.

Our Speakers:

- [Keegan Boyd](#), Partner, BLG
- [Don Boyer](#), President, BOYER@RegulatorySols
- [Edona Vila](#), Partner, BLG

Watch the webinar

By

[Keegan Boyd](#), [Edona C. Vila](#)

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BLG Offices**Calgary**

Centennial Place, East Tower
520 3rd Avenue S.W.
Calgary, AB, Canada
T2P 0R3

T 403.232.9500
F 403.266.1395

Ottawa

World Exchange Plaza
100 Queen Street
Ottawa, ON, Canada
K1P 1J9

T 613.237.5160
F 613.230.8842

Vancouver

1200 Waterfront Centre
200 Burrard Street
Vancouver, BC, Canada
V7X 1T2

T 604.687.5744
F 604.687.1415

Montréal

1000 De La Gauchetière Street West
Suite 900
Montréal, QC, Canada
H3B 5H4

T 514.954.2555
F 514.879.9015

Toronto

Bay Adelaide Centre, East Tower
22 Adelaide Street West
Toronto, ON, Canada
M5H 4E3

T 416.367.6000
F 416.367.6749

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