## Reviews

# **Canadian REB/IRB Review Services**

#### Localized Solutions Backed by Canada's Most Trusted Review Partner

Providing human subject protection oversight in Canada since 1993, Advarra's dedicated review panels in Ontario and Quebec expertly manage Canada-specific research.

Advarra has the most extensive site reach in Canada, and Canadian citizens comprise the majority of our Canadian review panel.

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#### **Regulatory Peace of Mind**

Strong understanding of Health Canada and U.S. FDA regulations as well as privacy requirements (e.g., HIPAA, PIPEDA)



#### Local Perspectives, Broad Experience

Strong understanding of current regulations and majority Canadian membership leveraging Advarra's diverse expert network



#### **Expertise for Every Trial**

Experienced in every therapeutic area and trial modality, including cell and gene therapies, Phase I, minimal risk, and community-based research

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#### **Technology-Enabled Efficiencies**

Real-time status details, optimized processes, and the most integrated IRB platform ensure approval documents are in order



#### French and English Support

Native French and English speakers and multilingual support for English, French-Canadian, and Spanish languages (including translation services and document verification)



#### Streamlined Process for North American Trials

Single submission for cross-border Canadian and U.S. studies and coordinated reviews to help ensure consistency and compliance



Advarra is the largest central IRB/REB in Canada that is AAHRPP accredited.

### **Turnaround Times**

\* Full Accreditation Research Protection Proposition

Advarra is committed to getting your trial started on time and helping you reach critical study milestones quickly.

| Review Item   | Submission to Decision |                   |
|---|------------------------|-------------------|
| New protocol and initial informed consent for multisite studies (full board review) |                        | 4-5 business days |
| New protocol and initial informed consent<br>(minimal risk review)                  |                        | 1-2 business day  |
| New site for a multisite study  |                        | 1 business day    |

Please note: Turnaround time is dependent upon complete and accurate submission of study documents. Any follow up requiring a response may extend this turnaround time.

#### Most Extensive Reach of Any Central REB/IRB

The only REB/IRB with official Canadian provincial recognition, able to service more of Canada than any other partner.

