

IBC-Ready™

Local IBC Submission Support

Institutional Biosafety Committee (IBC) submissions can be challenging for researchers new to the field of cell and gene therapy. Similarly, local IBCs may primarily focus on preclinical research review and might not be familiar with reviewing research involving human participants. This can lead to longer review timelines and a lot of back-and-forth clarifications, delaying patients' access to potentially life-altering research.

Advarra® Can Help Complete Local IBC Forms Accurately and Efficiently -Regardless of the IBC of Record

With IBC-Ready, Advarra can help reduce confusion and administrative burden by coordinating with the sponsor, site personnel and local IBC(s) to complete local IBC submission forms. Advarra packages all IBC documents as required by NIH quidelines and can submit the documents directly to the local IBC or to a site's study coordinator.

The IBC-Ready Advantage

Decrease timelines for local IBC submission and review

Reduce administrative burden for site(s), CRO and sponsor with a single entity coordinating submissions

Eliminate confusion for investigators and study coordinators unfamiliar with IBC questions

Service provided by biosafety professionals experienced in local IBC requirements and processes

Advarra's Streamlined Process for Local IBC Submissions Site A Site B Sponsor/CRO/SMO IBC (Site A) IBC (Site B) IBC (Site C)

Bettering Local IBC Submissions

Advarra's biosafety experts have extensive experience working with local IBCs and know how to answer IBC submission questions in a way that streamlines local review.

Even if Advarra® is not the site's IBC of record, our team will work with everyone involved to make sure the IBC submission forms are completed accurately and submitted in a timely manner so that the research can begin.

