Effective January 1, 2018, this charge replaces the previous fee structure for use of a commercial IRB (including WIRB)

To Whom It May Concern:

The institution has established a process by which certain types of industry sponsored research can utilize an AAHRPP accredited commercial Institutional Review Board designated by the sponsor. However, certain regulatory, administrative and other oversight responsibilities remain the responsibility of the University of Colorado Anschutz Medical Campus.

These responsibilities retained by the University include:
- Monitoring investigator training in human subject protects to assure it meets institutional and best practice standards;
- Ensuring that the necessary affiliated site specific requirements have been met;
- Acting as the privacy board for the protocol;
- Overseeing University and regulatory Financial Conflict of Interest policies and procedures;
- Providing regulatory compliance review when appropriate such as Institutional Biosafety
- Maintaining a compliance oversight role locally.

To off-set these administrative costs:
A one-time additional Research Regulatory Administration Fee of $5000 plus 28% F&A must be added to the clinical trial budget.

Yours sincerely,

Alison Lakin, RN, LLB, LLM, PhD
Associate Vice Chancellor for Regulatory Compliance