wcg IRB

EXPERIENCE MATTERS

For over 50 years, WCG IRB has built a legacy of trust, experience, and expertise as the oldest and most respected Central IRB in North America. Spanning the largest network of Central Sites & Institutions of any IRB, WCG IRB has reviewed over **62,000 studies** across nearly every therapeutic area in **over 300,000 sites** over the past **20 years** alone.

WCG IRB is the acknowledged industry leader in ethical and regulatory human research oversight– with multiple decades of IRB and IBC experience and trusted relationships with sponsors, CROs, institutions, and investigators.

93% IRB of record for 93% of all North American protocols.

92% Involved in 92% of FDA Approvals from 2019-2021.

20+ More than 20 successful FDA audits (dating back to 1983).

500+ 500+ successful Sponsor/CRO audits.

20x Holds IRB meetings up to 20x/week.

7 The first IRB in the industry to implement a 7-day work week/organization.

By leveraging MSA contracts with 3,300+ Institutions, AMCs, and Hospitals and ability to approve all private practice doctors, WCG can eliminate 45-60 days from IRB start-up timelines.

WCG IRB will help dramatically accelerate study start-up by significantly reducing turnaround time in the IRB review process and eliminating duplicative IRB review across Institutions and Private Practice sites.

WCG IRB will provide greater predictability and precision to site selection for studies through access to WCG's unique and comprehensive database of institutional information.

Due WCG IRB's industry-leading relationships with thousands of institutions, we relieve burdens for sponsor and CRO clinical operations teams by avoiding redundant local IRB reviews for any given study.

FDA ISO® AAHRPP Association for the Accreditation of Human Research Protection Programs, Inc.®