



# SITE CAPABILITIES PROFILE

Prepared for sponsor & CRO feasibility teams · June 2026

Independent multi-therapeutic research site · Three locations across Queens & Long Island, NY · Phase II–IV trials since 2013

**22,000+**

patient database, multi-ethnic

**Since 2013**

in clinical research

**5**

physician investigators

**Phase II–IV**

Phase I considered

## THERAPEUTIC EXPERIENCE

**Gastroenterology:** Crohn's disease, ulcerative colitis, IBS, chronic constipation, GERD

**Cardiology & Metabolic:** heart failure, hypertension, dyslipidemia, obesity & weight management

**Endocrinology:** Type 1 diabetes, Type 2 diabetes

**Neurology:** migraine, Alzheimer's disease & dementia

**Dermatology:** psoriasis, acne

**Respiratory:** asthma, COPD

**Rheumatology & Immunology:** rheumatoid arthritis, lupus (SLE), osteoarthritis

**Additional experience:** liver disease, urinary tract infections

*Specialties span internal & family medicine, cardiology, endocrinology, gastroenterology, pulmonology, neurology, psychiatry, rheumatology, and oncology — backed by a 22,000+ pre-consented patient database.*

## FACILITY & CAPABILITIES

- Three clinical locations — Merrick, Queens Village & Richmond Hill, NY
- Dedicated research space with private exam rooms
- Phlebotomy and on-site sample processing
- 12-lead ECG / EKG
- Refrigerated centrifuge
- –20°C and –70°C specimen storage
- IV infusion administration capability
- Up to 8 concurrent studies
- Equipment calibrated annually
- Private rooms for confidential informed consent
- Double-locked, temperature-monitored IP storage

*Imaging & procedures: dedicated offsite radiology (X-ray, MRI, ultrasound), biopsy, and at-home participant visits. Additional capabilities and services discussed to fit your protocol.*

## INVESTIGATOR BENCH

Five licensed physician investigators, each with more than twenty years of clinical and research experience, including dedicated specialists in cardiology and endocrinology. Our investigators conduct their research exclusively with A1 Clinical, and study oversight is performed by the investigators directly — never delegated down. Investigator CVs and current licensure are provided with feasibility responses.

## CONDUCT & COMPLIANCE

All studies conducted under ICH-GCP guidelines for global sponsors. Source documents and study materials are kept under secured, delegated access; the site meets HIPAA Privacy & Security Rule requirements. Feasibility questionnaires and regulatory documents are turned around promptly.

### Start a feasibility conversation

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