Remington-Davis Clinical Research

Remington-Davis Phase 1 Capabilities & Experience

Remington-Davis, Inc. (RDI) is an independent clinical research site with more than 30 years of continuous operations and over 600 completed clinical trials. We conduct Phase I–IV trials and have strong, proven capabilities in early-phase research, including healthy volunteer, vaccine, and specialty-population studies.

Phase 1 Infrastructure & Operational Capabilities

- Five private beds for overnight confinement.
- Outpatient Phase 1 capacity with confined dosing and PK/PD sampling.
- Private exam rooms and dedicated infusion/sampling rooms.
- On-site laboratory processing with access to partnering local lab.
- Secure temperature-controlled investigational product storage. Limited access and continuously monitored. Including DEA scheduled drugs.
- 12-lead EKG machines and pulmonary function testing room.
- 24-hour coordinator coverage during confinement periods.
- Subject lounge and observation areas for continuous monitoring.
- On-site shower and kitchenette.

Recent Phase 1 Experience (Last Three Years)

- Healthy Volunteer Phase 1 Vaccine (EBV): 34 participants enrolled to date; ongoing.
- Older Adult Phase 1 Study (ages 65+): 85 participants enrolled; normal-coagulation population.
- Capabilities demonstrated: PK sampling, confined dosing, continuous safety monitoring.

Broader Phase 1 Experience

- Biologic therapy: first-in-human anti-coagulation reversal agent for ticagrelor (Brilinta).
- Phase 1 vaccine studies in Escherichia coli, Streptococcus, Clostridium difficile and Epstein-Barr virus.
- Extensive early-phase infectious disease and immunology experience.
- Atopic Dermatitis Phase 1
- Psoriasis Phase 1 Study

Participant Profiles & Recruitment Strength

- Healthy adults: located 2 miles from The Ohio State University, providing access to large, diverse young-adult populations ideal for Phase 1 research.
- Specialty populations: strong access to older adults (including 65+), metabolic disease, cardiovascular risk groups, and other internal-medicine categories.
- Internal database of more than 20,900 participants with approximately 37% diversity.
- Full-time recruiters and targeted outreach across Central Ohio.

Why Sponsors Select Remington-Davis for Phase 1

- Rapid start-up: contract and budget turnaround typically within 3-5 business days.
- Central IRB friendly with fast regulatory submission.
- High retention and compliance supported through concierge-level participant services.
- Experienced coordinators accustomed to early-phase operational intensity.
- Independent site: flexible, responsive, and highly sponsor-focused.

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