REMINGTON-DAVIS
Professional Clinical Research

CASE STUDY

Accelerated Patient Recruitment During Pandemic Conditions

OVERVIEW
Who We Are

Remington-Davis is an independent clinical research site located in Columbus, Ohio, with extensive experience supporting sponsors and CROs across multiple therapeutic areas. This case study highlights our ability to rapidly activate and execute a study under compressed timelines and challenging external conditions.

CHALLENGE
Study Background

Remington-Davis was approached by a medically focused Contract Research Organization (CRO) to provide site services for a migraine glasses study. The study experienced initial regulatory setbacks, having been rejected by the FDA before later receiving approval.

Following FDA approval, the sponsor faced significant pressure to accelerate study timelines, creating an urgent need for rapid site activation and patient recruitment.

STRATEGY
Key Challenges

Several factors contributed to the complexity of study:

Compressed Start-Up Timeline: Due to the FDA approval delay, timelines were significantly shortened, requiring expedited site readiness and activation.

Pandemic Conditions: The study was conducted during the COVID-19 pandemic, introducing challenges related to patient access, safety protocols, and operational logistics.

Patient Recruitment Barriers: Public health concerns and restrictions made traditional recruitment strategies less effective.

RESULTS
Remington-Davis Approach

To overcome these challenges, Remington-Davis implemented a focused and adaptive strategy:

Rapid Study Activation: Streamlined internal processes enabled quick regulatory and operational readiness.

Targeted Recruitment Strategy: Leveraged existing patient databases, digital outreach, and community engagement to identify qualified participants efficiently.

Flexible Patient Engagement: Implemented safety-conscious scheduling and communication protocols to maintain patient confidence and participation. Operational Agility: Adjusted workflows in real time to address evolving pandemic-related constraints.

60 PATIENTS ENROLLED
In 60 days

MAINTAINED HIGH LEVELS
of patient engagement and retention

MET SPONSOR EXPECTATIONS
for speed and quality

CONCLUSION
Why It Works

This study demonstrates Remington-Davis' ability to execute under pressure, adapt to rapidly changing conditions, and deliver strong enrollment outcomes. Our proactive approach, operational efficiency, and patient-centric strategies make us a reliable partner for CROs and sponsors seeking high-performing clinical sites.

For additional information about our capabilities, we welcome the opportunity to connect.

Every breakthrough has a beginning.
Start yours.

READY TO TALK?
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