



REVIVAL
RESEARCH INSTITUTE, LLC

ADVANCING RESEARCH, EMPOWERING SITES

YOUR TRUSTED CLINICAL RESEARCH PARTNER


EXIT
PRIME
REVIVAL

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Company Overview

Revival Research Institute is a multi-disciplinary clinical research organization dedicated to simplifying clinical trials for sponsors and improving access for patients. From feasibility and start-up to recruitment, data, and close-out, we manage every step under one roof.

Mission

To bring research closer to communities while delivering trials that are faster, reliable, and patient-centered.

Vision

To set the benchmark for community-driven clinical research, where every trial is efficient, inclusive, and trusted.



Revival's Operational Footprints

Michigan, Texas, Nebraska, New Jersey, North Carolina, Illinois, & Poland

25+ sites
Interspersed Throughout The States



30+ Principal Investigators & **27+** Sub-Investigators throughout the States

Over **200** studies conducted yearly in over **12+** therapeutic areas throughout the States

Pillars That Define Us

Running a clinical trial shouldn't feel like navigating a maze of delays, dropouts, and endless data questions. Our commitment is built on three core pillars fast study start-up, highest patient recruitment, and query free data that ensures every study launches faster, enrolls better, and delivers data you can trust with confidence.

Fast Study Start-up

We activate sites quickly speeding contracts, ethics approvals, and site initiations so trials launch without delay.

Highest Patient Recruitment

Our strategies deliver high enrollment and diversity, while strong retention keeps participants engaged to study completion.

Query-Free Data

Meticulous QC and standardized systems ensure accuracy, fewer queries, and smoother regulatory approvals.

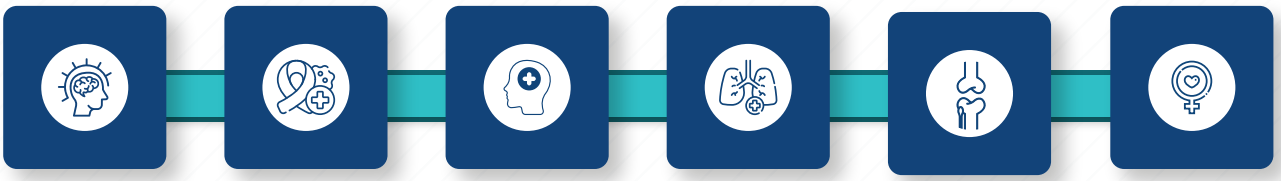
200+
Studies Conducted Annually

82%
Met/Exceeded Enrollment Target

95%
Patient Retention Rate

Areas Of
Therapeutic Expertise








LEARN M
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REVIVAL
INCREASING ACCESSIBILITY OF
REVIVAL




REVIVAL'S
SITE NETWORK
REVIVAL

Our Strengths

Established Industry Connections

- Strong Relationships with Sponsors & CROs
- Access to Top-of-the-Line Technology Systems
- Collaborative Partnerships with Leading Sites
- Continuous Flow of High-Quality Studies





Proven Recruitment Engine

- Dedicated Patient Recruitment Teams
- Nationwide Pharmacy Partnerships

- Referral & Transport Programs
- Pre-Screening Initiatives for Higher Enrollment

Rigorous Trainings

- Quarterly All-Hands Trainings
- Extensive Online Training Courses

- Bi-Weekly Compliance Trainings & Check-ins
- Skill/Role-Based Training Programs



Competitive Edge

Industry Standard

- **SITE INITIATION TIME**
  2 months
- **DATA UPLOAD ON EDC**
  3-5 days
- **REGULATORY DOCUMENTS UPLOAD**
  5-7 days
- **FOLLOW-UP ON PATIENT LEADS**
  3-5 days

Our Timelines

- **SITE INITIATION TIME**
  2 weeks
- **DATA UPLOAD ON EDC**
  Within 24 hours
- **REGULATORY DOCUMENTS UPLOAD**
  Within 48 hours
- **FOLLOW-UP ON PATIENT LEADS**
  1 hour

COMMITTED TO REDEFINING PATIENT CARE

ONE CLINICAL TRIAL AT A TIME



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