



# InoRx

Simplifying Trials. Accelerating Cures.



## Corporate Fact Sheet

### About InoRx

InoRx is a next-generation CRO built to accelerate clinical development through innovation, patient access, and operational excellence. Headquartered in the U.S. with operations in India, InoRx leverages its affiliate network of academic medical centers and clinics, high quality standards, and experienced clinical operations, to deliver faster, more cost-effective, highly reliable clinical studies for sponsors worldwide.

### Backed by SRM Group

InoRx is part of the SRM Group, a diversified global enterprise with \$1.4 billion in revenue and 10,000+ employees across healthcare, education, technology, and more. SRM's decades of experience and reputation in medical research, hospital systems, and academic excellence provide InoRx with scientific expertise, infrastructure, and access to a vast network of clinicians, research sites, and patients.

### Therapeutic Focus Areas

- Cardiovascular
- Oncology
- Endocrinology/Metabolic Syndrome/Nephrology (Diabetes, Obesity, NASH)
- Virology/Infectious Disease/Vaccines
- Respiratory/Pulmonary Disease



## Core Capabilities



**Phase I Clinical Trial Capabilities:** ICMR-recognized early-phase unit with specialized infrastructure and experienced investigators delivering first-in-human, SAD/MAD, vaccine, and device studies with rapid start-up and robust safety oversight



**Patient Centric Trial Design:** Design smarter protocols by uniting deep therapeutic expertise with patient insights to drive faster enrollment, higher retention, and accelerated approvals



**Feasibility & Enrollment Planning:** Combine real-world data, physician insights, and predictive modeling to identify patients with precision and accelerate recruitment



**Accelerated Site Start-up:** Streamlined regulatory and contractual workflows to launch trials in weeks, not months



**Medical Writing:** Expert protocol development and study reporting with scientific precision



**Execution Services:** End-to-end project management ensuring compliance, patient safety, faster patient enrollment, and on-time delivery



**Data Management & Biostatistics:** Deliver clean, compliant data and rapid, reliable analyses that accelerate decisions and drive regulatory success



**Pharmacovigilance & Clinical Trial Safety Services:** End-to-end safety management including SAE/SUSAR processing, medical review, expedited reporting, safety surveillance, and regulatory submissions to ensure patient safety and protocol compliance



**Laboratory Services:** In-house central and specialty lab capabilities with standardized processes that ensure sample integrity and regulatory compliance



**Site Network Solutions:** SRM hospitals, 150+ rigorously vetted sites, 800+ Primary Investigators, and large patient populations across diverse therapeutic areas to accelerate enrollment

## What Sets InoRx Apart

### Accelerated Patient Access:

Direct access to SRM's extensive hospital network unlocking a population of 500,000 across key therapeutic areas under a unified governance structure enabling faster recruitment and earlier proof-of-concept.

### Tailored to Your Outcomes:

InoRx is purpose-built for small and mid-size sponsors, providing senior leadership attention, flexible engagement, and tailored solutions that align with the urgency of biotech development.

### Trusted Foundation in Care

**& Research:** Backed by SRM's reputation in healthcare and academics, InoRx brings credibility, stability, and experienced clinicians to ensure trials are conducted to global standards with uncompromising quality.

