



Bringing safe, effective, and  
affordable therapies to patients faster

## The costs of delayed and prolonged clinical trials can be very high!

Efficient drug development cycles are extremely crucial to ensure patients are provided with timely access to safe and effective therapies. Though, drug development is a complex process and can take a minimum of 10-15 years from product identification to commercialization.

Clinical trials, if managed right, can help bring down the arduous timelines of drug development, bring therapies to patients faster, and save millions of dollars for pharmaceutical companies. Trial participant experiences heavily influence enrollment and retention, and therefore successful completion of trials.



### Clinical development teams face multiple challenges in trials

- **Increasing complexity of clinical trials**  
Rapid adoption of decentralized and hybrid trial formats, as well as precision therapies
- **Delays in clinical study startups**  
Lack of robust processes and insights for site selection, activation, screening, and enrollment
- **Poor patient retention levels**  
Sub-optimal use of technology and RWD/RWE towards improving patient experiences

### We simplify clinical development lifecycles, to drive faster decisions and predictable success

- **Proven trial acceleration solutions**  
Reproducible roadmap to effective adoption of digital
- **End-to-end management of trial operations**  
Accountability from trial design to completion
- **Risk mitigation and scalability**  
Deep domain expertise to inform strategy and drive actions

### Here's how we can help you

#### Bold Vision

build capabilities, culture

**Future Ready  
Healthcare**

Healthcare expertise,  
content and conversations

#### Accelerate

and scale up outcomes

**EXT**

Fit-for-purpose  
technology

#### Personalize

experience

**indegene**

Agile operations  
and culture

## Technology-led solutions that span across the clinical R&D lifecycle including study startup, conduct, and closeout

Feasibility Assessment	Startup	Conduct	Closeout
<ul style="list-style-type: none"> <li>• Trial landscape and competitive intel</li> <li>• Site selection strategy</li> <li>• Trial design</li> <li>• Trial disclosures</li> <li>• Regulatory strategy</li> <li>• Recruitment forecasting</li> </ul>	<ul style="list-style-type: none"> <li>• Site selection</li> <li>• Site initiation</li> <li>• Patient screening</li> <li>• Patient enrollment</li> <li>• Digital study portals with IRB approved content</li> <li>• Accelerated study startup, budgeting, contracting, and payments</li> </ul>	<ul style="list-style-type: none"> <li>• Digital site and patient engagement</li> <li>• Site monitoring                             <ul style="list-style-type: none"> <li>• Risk-based quality management (RBQM)</li> <li>• Medical monitoring and review</li> </ul> </li> <li>• Data management, programming, and standardization</li> <li>• Clinical data integration and aggregation</li> </ul>	<ul style="list-style-type: none"> <li>• Document and submission publishing</li> <li>• Study documentation and content handling</li> </ul>
E2E program and project management with complete oversight and transparency			

### Here's how we are different

#### AI/ML-driven trial accelerator solutions

- Trial designs customized to disease landscape, patient heterogeneity, and expected outcomes
- Integrated platforms, with the ability to generate clinical trial data, early insights, and evidence for informed decision-making

#### Risk-based quality management (RBQM)

- Real-time dashboards with visibility to track trial progress and associated risks while triggering timely actions

#### Patient-centric trial ecosystem

- RWD/RWE to refine patient experiences
- Multi-site presence across geographies and ethnicities
- Wide coverage of prequalified digital site networks

#### Deep domain expertise

- Multidisciplinary team of Data Analysts, Statisticians, Data Scientists & Engineers, Clinical & Medical SMEs

### Here are just a few outcomes we delivered

>75%	>800	40%	65%	<30 min
Reduction in time for preliminary feasibility	Clinical Study Feasibility Analyses	Faster study startup	Wider reach on eligible patient population	Response time with 24x7 concierge services
30%	>95%	20%	1000+	
Reduced risk of study failures	Patient retention for completed studies	Reduced disclosure and submission time	Study migrations	

Through local teams, we support healthcare organizations  
wherever they are



## About Indegene

We are a technology-led healthcare solutions provider. We combine deep industry expertise with fit-for-purpose technology in an agile and scalable operating model. Many of the leading global healthcare organizations rely on us to deliver effective and efficient clinical, medical, and commercial outcomes every day. From strategy to execution, we enable healthcare organizations be future ready.

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