

End-to-End Confidence: Intelligent Trial Designs, Trusted Data, Expert Delivery

+30%

Probability of Success

-50%

Development Costs

-40%

Trial Duration & Required Enrollment

150+

Global Customers

700+

Studies

25+

Submission Packages

PhaseV Advantage

Smarter, Faster Trials

- AI-Powered Optimizer – Optimizes designs and enables real-time adjustments
- Predictive Modeling – Forecasts enrollment, retention, and endpoints
- Dynamic Decision Support – Adjusts trials for faster, cost-efficient execution
- System Integration – Connects securely with EDC, CTMS, and workflows

Bioforum Advantage

Reliable, Compliant Execution

- Data Management – Accurate, end-to-end data lifecycle management
- Biostatistics & Programming – CDISC-compliant, advanced modeling
- Operational Expertise – Efficient global trial execution
- Regulatory Alignment – FDA, EMA & ICH-compliant delivery

➤ Accelerated Timelines

Faster, more efficient trials with optimized designs and expert execution

➤ Cost Efficiency

Lower costs through fewer amendments and streamlined operations

➤ Trusted Data & Insights

Trusted, submission-ready data from predictive analytics

➤ Scalable Across Phases

Applicable to all phases (I–IV) and therapeutic areas with oncology expertise

