

Global regulatory strategy, local expertise

Grow your product's potential worldwide

You've already invested so much in your life-changing therapy. Don't leave its success to chance. Lean on deep regulatory expertise throughout your product's lifecycle to manage global complexity, and strengthen your market approach.

Navigate the regulatory approval process

Stay ahead of constantly changing regulatory requirements across multiple regions with local market expertise

Minimize resource expenditure and the need for your internal resources and workstream management

Boost your chances of more efficient launches and earlier revenue streams

Manage global complexity

Quickly identify and address regulatory/compliance risks through deep understanding of the local market

Streamline operations and reach further with our worldwide network of on-the-ground expertise

Expand your regulatory capabilities

Enhance flexibility in meeting global-to-local needs, bolstering your market presence and supporting your expansion strategy

Reduce costs by leveraging our existing infrastructure, including everything from local experts to project managers

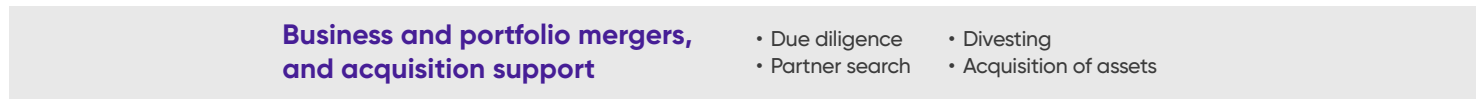
From pre-clinical to commercial, we strive to accelerate speed to market



Integrated product development ———— Innovations to market ———— Optimal patient access



←————— **Strategy and scientific consultancy** —————→



Acronyms

BLA	Biologics license application	HTA	Health technology assessment
CMC	Chemistry, manufacturing, and controls	IND	Investigational new drug
CRO	Contract research organization	MAA	Marketing authorization application
EAP	Early access program	PV	Pharmacovigilance
GxP	Good practice guidelines and regulations	QMC	Quality, management, and compliance
GPO	Group purchasing organization	RWE	Real world evidence
HEOR	Health economics outcomes research	TPP	Target product profile