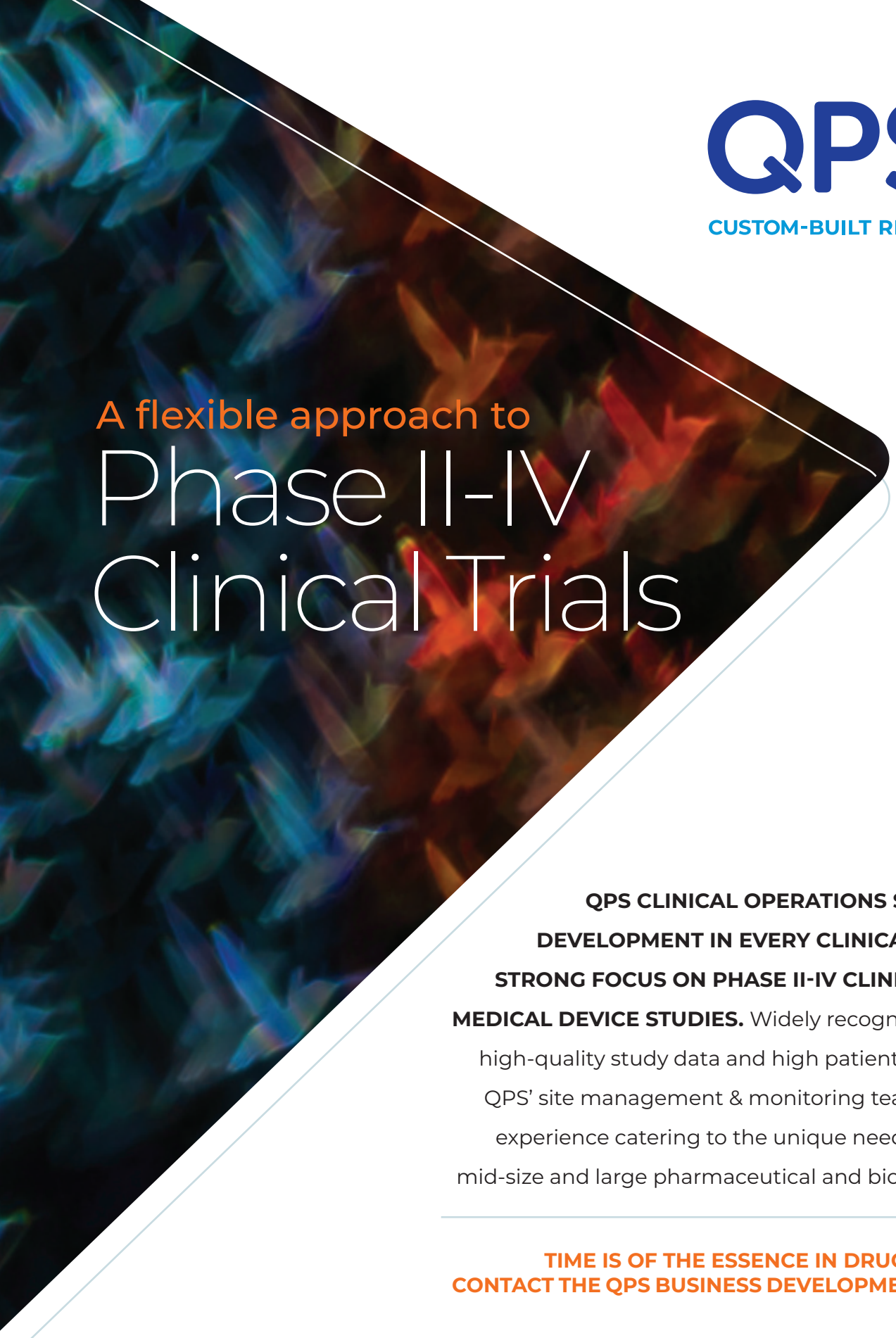

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QPS CLINICAL OPERATIONS SUPPORTS DRUG DEVELOPMENT IN EVERY CLINICAL STAGE WITH A STRONG FOCUS ON PHASE II-IV CLINICAL TRIALS AND MEDICAL DEVICE STUDIES. Widely recognized for achieving high-quality study data and high patient enrollment rates, QPS' site management & monitoring teams bring years of experience catering to the unique needs of virtual, small, mid-size and large pharmaceutical and biotechnology firms.

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Therapeutic Expertise Includes:

QPS has extensive experience across multiple Therapeutic Areas, including: Neurology, Pain, Inflammation, Vaccines, Dermatology, Gynecology, Urology, Diabetes, Metabolic Disorders, Infectious Disease, CNS, Respiratory, and Rare and Orphan Diseases.

Key Differentiators:

We combine scientific leadership with disciplined execution, global access to patients, and local and global regulatory expertise to accelerate your compound to approval.

- Full-time physicians provide strategic direction for study design and planning, train operations staff, work with primary investigators, provide medical monitoring, and meet with regulatory agencies as needed.
- Long-standing interactions at the medical and operational levels with global key academic and industry opinion leaders.
- Close collaborations and ongoing programs with high recruiting sites at academic, network, and independent centers.
- Our expertise as a site management organization and a full service late phase CRO provides us instrumental insight into the needs and challenges of the sites allowing us to proactively manage and support them.
- Our unique operational platform allows us to cost effectively place the most experienced and therapeutically focused team members on your project.
- A proven study initiation process that supports and encourages team work, transparent communication, risk mitigation insight and individual ownership of expectations.

Professional and Experienced Staff

With QPS you get immediate access to our Global Clinical Development Teams who are capable of meticulously managing the conduct of your trial while working collaboratively with your development team.

From the first interaction we are there to provide insight, consultative services, protocol review, site feasibility and recommendations. We will evaluate your protocol to determine the right project team mix and experience necessary to successfully perform your trial within the designated time lines and within budget.

Whether the trial is to be performed in one country or many, we have the regional coverage, regulatory prowess and medical support to ensure a smooth start-up and selection of the right sites. With offices throughout the world we are in constant contact with thought leaders and regulatory authorities to help ensure the success of your trial.

Our project management team offers a diverse therapeutic expertise as well as years of management experience. If there should be a need for a specific indication that falls outside our in-house experience we will, in full transparency, offer up consultants that have that expertise.

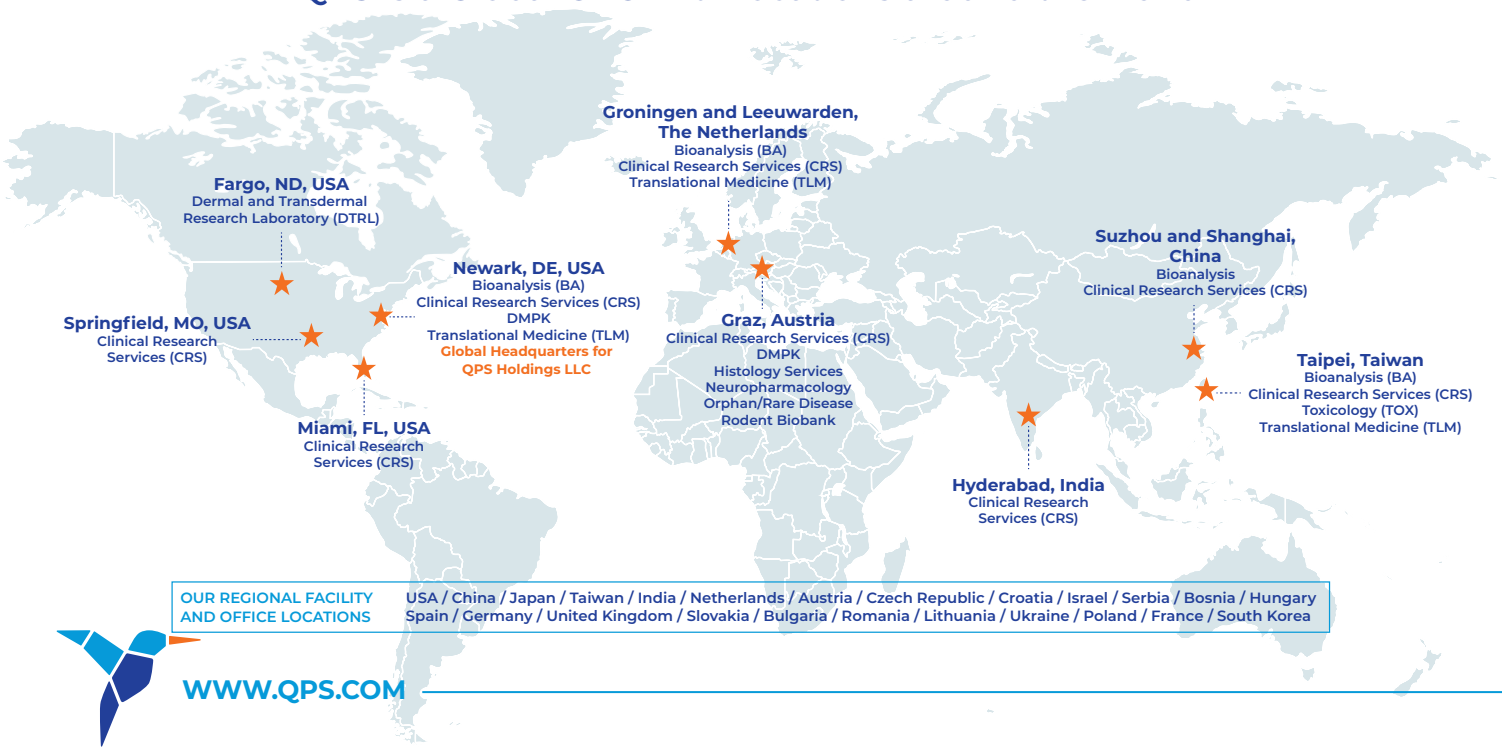
We believe you come to CROs for their experience, operational depth, in-house consultation, and geographic reach. We work hard to ensure that our clients are impressed with our teams on all levels which helps establish long lasting relationships.

"Please give QPS a call to see how we can work together to establish a great relationship of trust and mutual respect."

Kevin Vernarec

Sr. VP, Global Head of Late Phase Clinical Development

QPS is a Global CRO with locations around the world



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