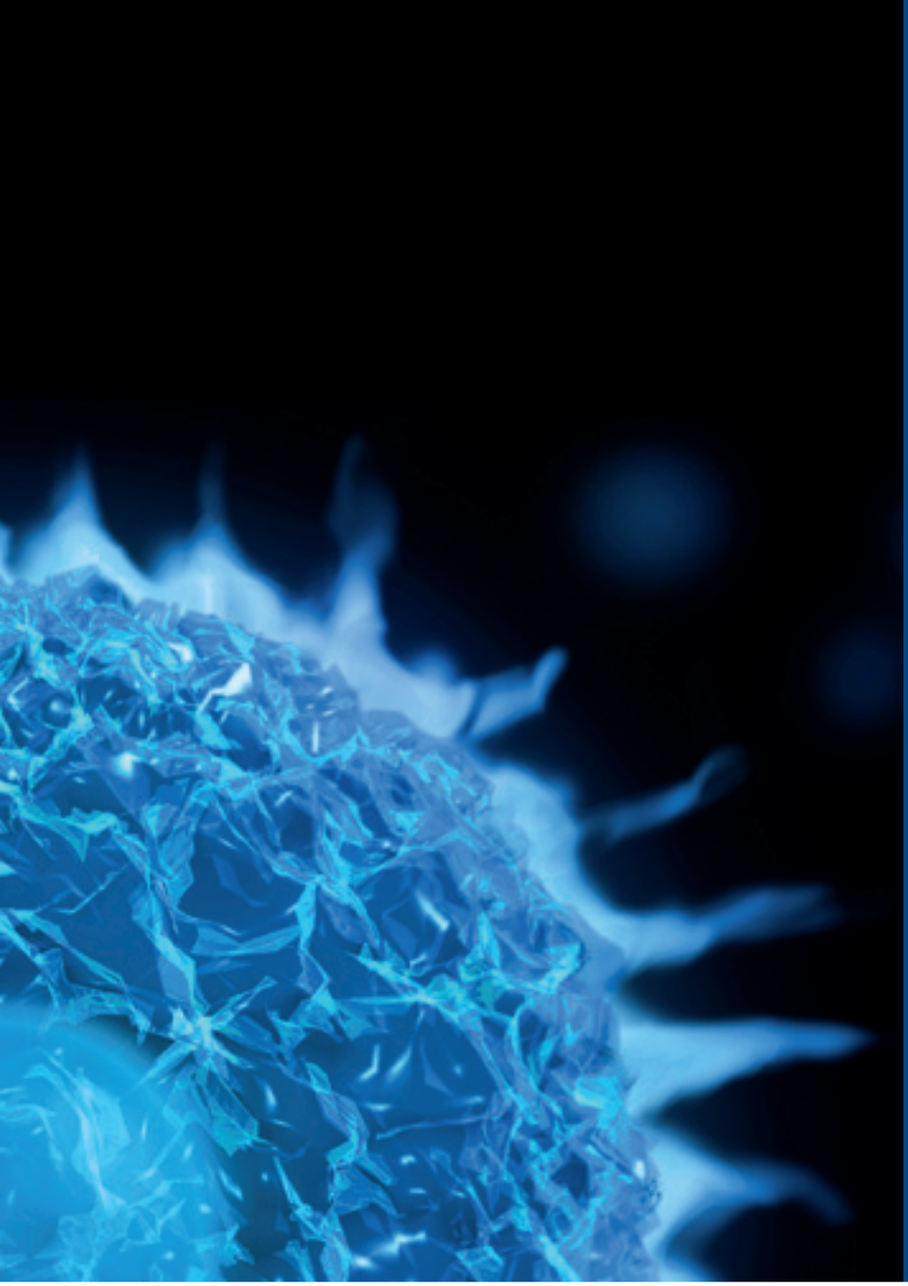
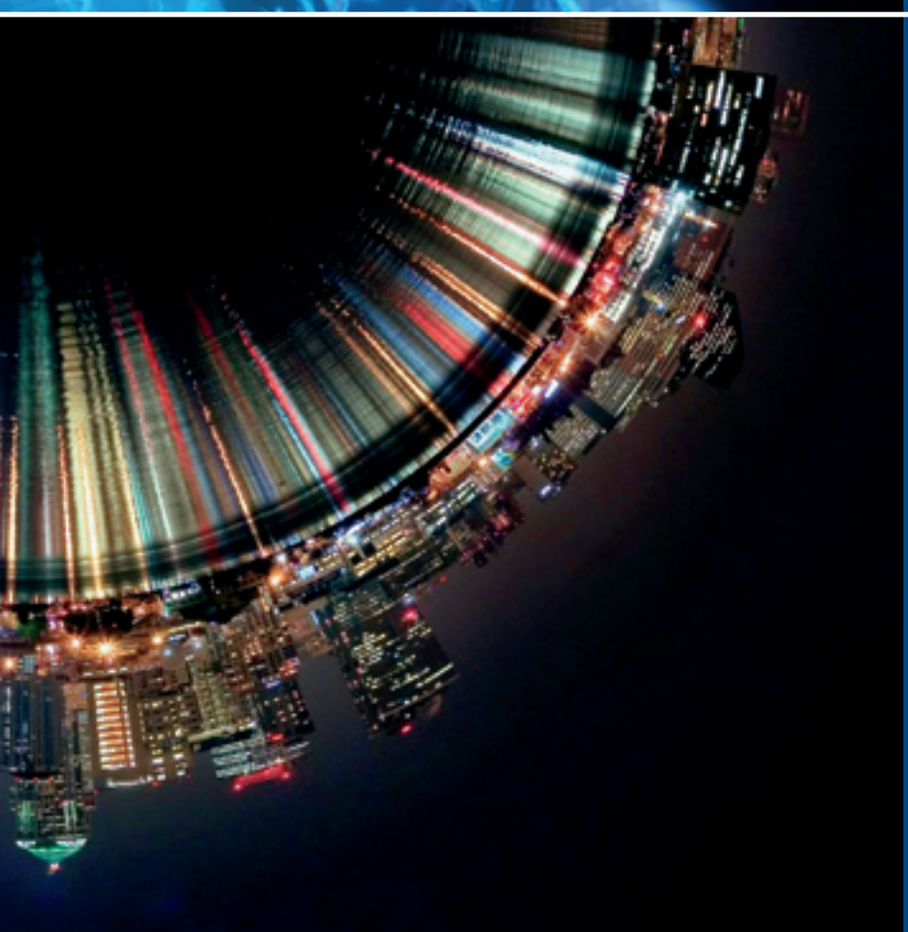


READYING YOUR SCIENCE FOR THE REAL WORLD



PAREXEL®

YOUR JOURNEY. OUR MISSION.™

Drug development today is more than a process. It's a journey.
At PAREXEL, we are committed to making your journey our mission. To solving problems before they arise. To making the process smarter and more efficient. To being a true partner to our clients, in every sense. For over 30 years we have helped our clients get their new and innovative drug treatments into the hands that need them most by simplifying their journey to market. Today that means going above and beyond—at every step along the way.

PAREXEL®

NAVIGATING THE JOURNEY TO MARKET TODAY REQUIRES A SPECIAL TYPE OF PARTNER.

At PAREXEL we bring together the most talented minds, operational excellence and technology in ways no one else can match. The results benefit our clients in three ways:

INTEGRATION

A GLOBAL MIND

Global harmonization allows our teams across the world to serve you effectively as one.

INNOVATION

DRIVEN TO SOLVE THE COMPLEX

Pervasive innovation and continual optimization help simplify the journey at every step.

PARTNERSHIP

THE BIGGER JOURNEY

A culture of deep collaboration helps accelerate development cycles and reduce costs.

FROM GLOBAL FOOTPRINT TO GLOBAL MIND.

While no two journeys are exactly alike, our clients share similar desires to access diverse patient populations, reduce study costs and conduct high-quality clinical research worldwide. Today that requires intelligence that only a fully integrated global network can provide.

Helped develop

95% of the

200 TOP-SELLING

biopharmaceuticals on the market

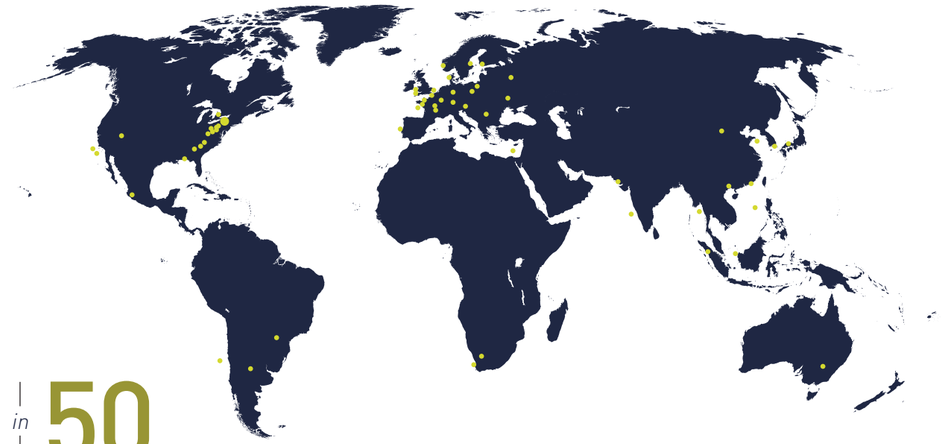
625

REGULATORY PROFESSIONALS

on staff

A SINGLE

SHARED TECHNOLOGY PLATFORM



76 OFFICES *in* **50** COUNTRIES

and growing

We currently support over

1,700

CLINICAL PROJECTS

in

20 THERAPEUTIC AREAS

We are 76 offices across 50 countries and still growing. Our offices are staffed by experts with local knowledge—experts who are prepared to advise on everything from the right patients to the right investigators and sites for your needs. They are also acutely aware of local regulatory requirements, so you can address any potential challenges early in the process. But what truly sets us apart is how we enable the teams across our global network to support you as one. We have invested significantly in global harmonization, which means we ensure your studies are conducted to the same standards of excellence, from Boston to Beijing.

Strategic Partnerships enable

DEEP **COLLABORATION**

We work with all of the **top**

50 **LARGE** biopharmaceutical companies

and all of the **top**

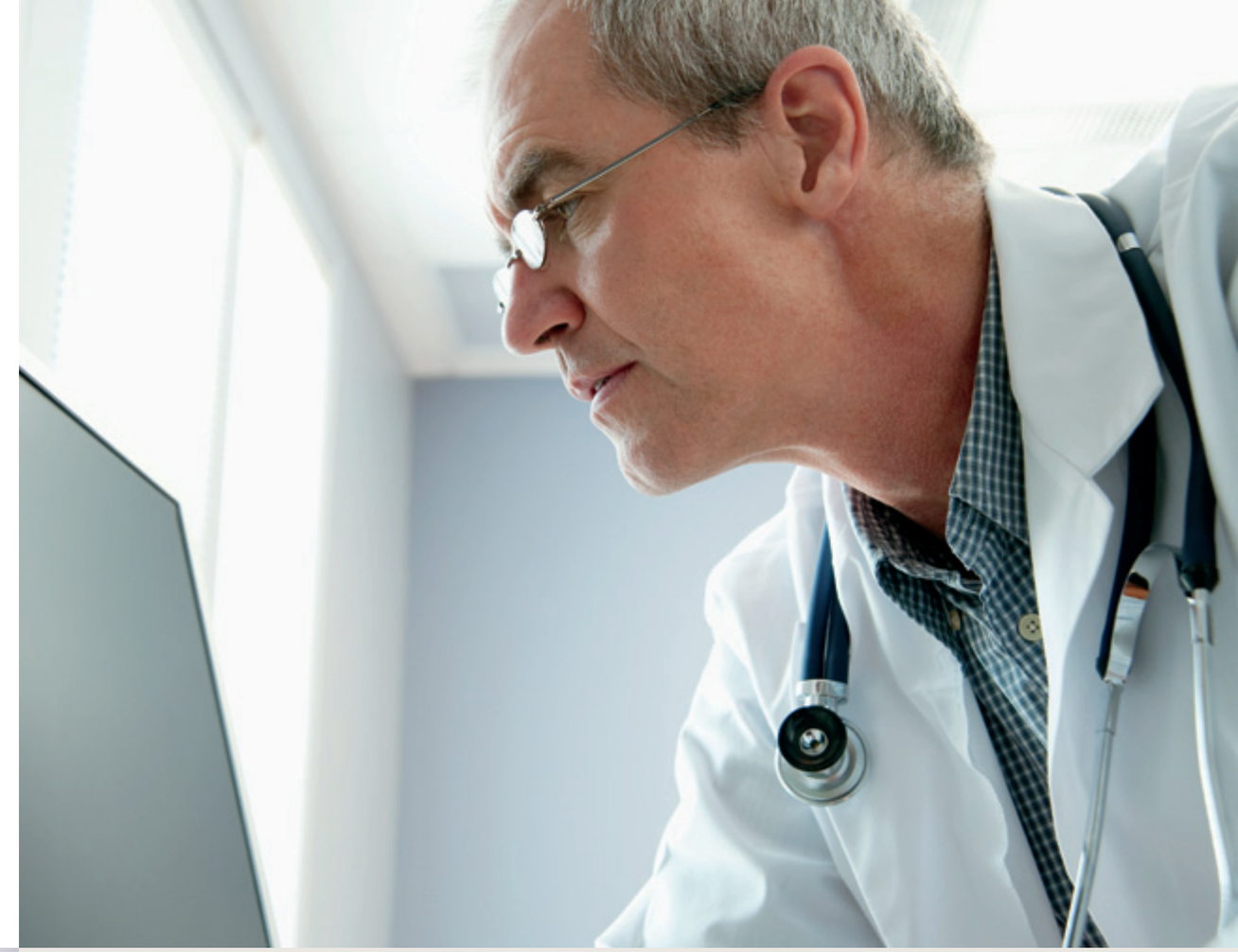
30 **SMALL & EMERGING** biopharmaceutical companies

We can conduct trials in more than

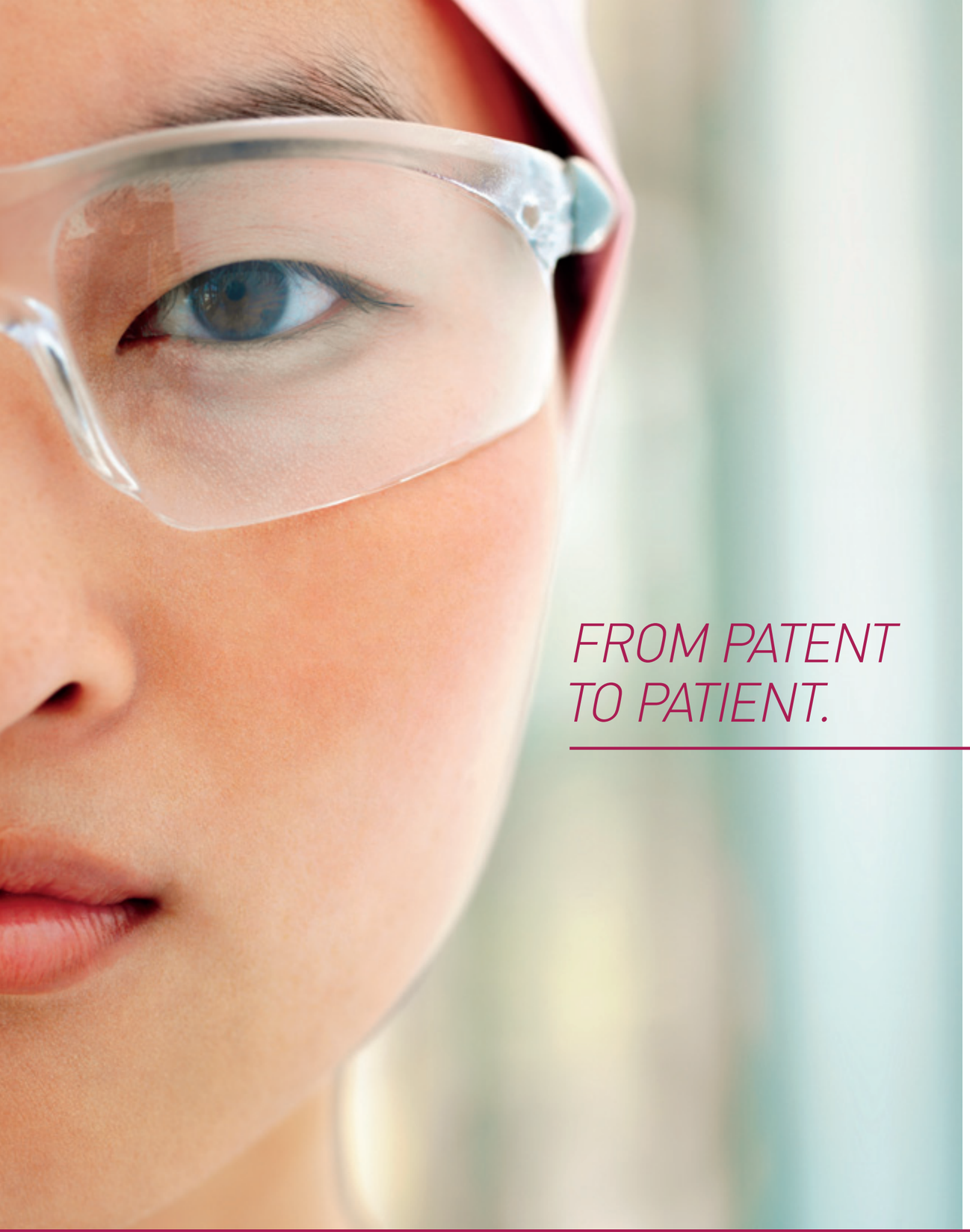
100

COUNTRIES

EVERY STEP OF THE JOURNEY IS AN OPPORTUNITY TO WORK SMARTER.



Though we are well known for our clinical research services, our breadth of expertise encompasses the continuum of product development and commercialization, from early planning through regulatory and post-approval. We view each step as an opportunity to work smarter, more efficiently and more effectively. We don't fear complexity, we embrace it. This mindset, backed up by achievement, has also enabled us to do pioneering work in emerging fields like biosimilars. Work that is paving the way forward for our clients and the industry.



FROM PATENT TO PATIENT.

PLANNING

The best-laid plans have contingencies built in.

Clinical trials management is only half of our story. We are equally strong in planning clinical development. Our consulting offerings cover the entire continuum of product development and commercialization. We bring upfront strategic thinking to the table, with an eye on your end objective, considering everything from the pathway you want to take, to the market, to your regulatory strategy, and even to the fit with payers.

THE CLINICAL PATHWAY

Early Phase

Each clinical trial phase requires increased investment, but the early phases are the foundation on which everything else builds. PAREXEL specializes in Early Product Development. We also have dedicated Early Phase facilities in five global centers and all of the expertise required to create cost-efficient development strategies that help our clients make better early-stage decisions.

Mid-Late Phases

As the scope of each study increases, so do the challenges. PAREXEL rises to meet them with the largest integrated patient recruiting capability in the industry, logistics and technology support that are second to none, and a global network that works seamlessly together to make sure your global study executions are as smart as your plans.

Peri-Approval

Peri-approval studies are on the rise. Effective studies can assist not only in fulfilling regulatory requirements and gaining market approval, they can also help maximize return on investment by bridging drug development and commercialization. PAREXEL supports clients with systems and strategies specifically developed for Phase IIIb studies that can ultimately help extend product lifecycle, expand prescribing communities and increase product exposure.

POST-APPROVAL

A new starting line.

Regulatory approval is not the end of the journey. The marketplace today demands the demonstration of real-world value before fully adopting a new treatment alternative. In this environment, late-phase observational research has emerged as a key tool to monitor safety and demonstrate both comparative and cost effectiveness.

PAREXEL combines deep experience on some of the largest Phase IV studies ever conducted with integrated technology that streamlines study operations to provide leading-edge service in this massively complex—and increasingly critical—arena.

WORKING SMARTER BY WORKING TOGETHER.

PAREXEL is committed to creating partnerships that deliver value, drive performance and bring new drugs to market faster. We continue to develop new models that blend client and service-provider resources for maximum efficiency. We are also redefining industry standards for service and what one can expect to gain from the right partnership. Whatever your needs, PAREXEL is committed to delivering solutions that make sense and to forming a productive relationship that works for you.

STRATEGIC PARTNERSHIPS

A shift in the industry.

As the needs of our customers have evolved, so has PAREXEL. Increasingly, biopharmaceutical companies are moving away from tactical engagements with us in favor of longer Strategic Partnerships, which are designed to deliver greater benefits throughout the development lifecycle. Strategic Partnerships can help accelerate development cycles while significantly reducing overall costs. A Strategic Partnership with PAREXEL is characterized by a deeper sense of partnership and a mutual investment in the relationship, as well as the inclusion of longer-term planning, which together help our clients achieve efficiencies across the board.

Our Strategic Partners are able to:

- Leverage overhead and infrastructure
- Complement internal knowledge with external expertise
- Improve the predictability of their R&D efforts
- Rapidly expand global reach
- Increase operational efficiencies
- Streamline the development process

FUNCTIONAL SERVICES

A la carte solutions that complete your picture.

We don't believe in one-size-fits-all solutions when it comes to our working relationships. Our Functional Services unit enables clients of all sizes to take advantage of our world-class support for specific functions—without committing to a full research program. In addition to getting just what they need, clients who leverage this model continue to benefit from our continuous process improvements and innovations.

PAREXEL BIOPHARM UNIT

A dedicated unit especially for small and emerging biopharmaceutical companies.

When the stakes are high, the PAREXEL BioPharm Unit makes sure you are always the priority. We assign a dedicated team, including a proactive senior management leader who will identify and mobilize the right resources from across our organization. We focus on delivering innovative and effective study execution, based on aligned incentives, so you can be confident about meeting each critical development milestone.

PAREXEL RECEIVES SOME OF THE **HIGHEST AND MOST CONSISTENT RATINGS** IN THE INDUSTRY.*

**Source: Industry Standard Research (ISR)*

A NETWORK OF SERVICES. AN INTEGRATED APPROACH.

Today, the top 15 biopharmaceutical companies use our technology solutions. The secret to our success, as with all of our services, lies in our integrated approach. In addition to retaining over 3,000 dedicated technology and informatics professionals, our technology solution designs are informed by experts from across our organization, including Clinical Research Services and Consulting. Experts who have decades of experience, and who understand what's really needed at each step of the journey.

PAREXEL INFORMATICS

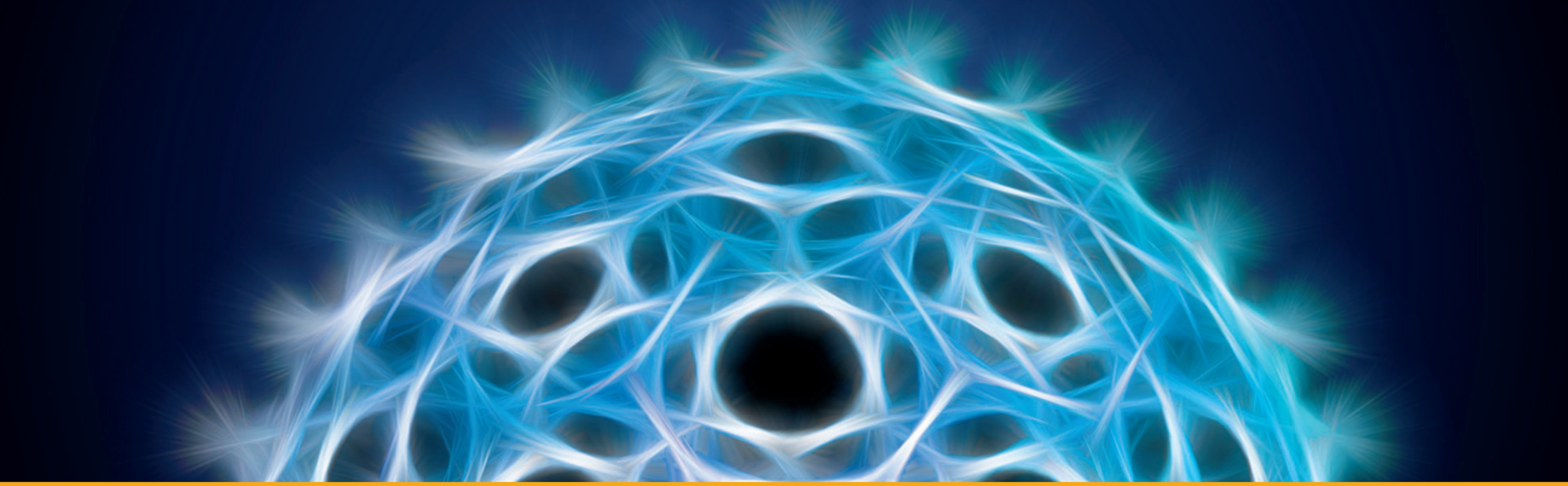
- Perceptive MyTrials® Platform & Infrastructure
- Study Management and Monitoring (IMPACT® CTMS)
- Clinical Data Management (DataLabs® EDC)
- Randomization and Trial Supply Management (ClinPhone® RTSM)
- Electronic Patient-Reported Outcomes
- Medical Imaging
- Regulatory Information Management (LIQUENT InSight,® LIQUENT SmartDesk™)

PAREXEL CLINICAL RESEARCH SERVICES

- Phase I: First-in-Human
- Phase IIa: Proof-of-Concept
- Phase II–III: Pivotal Clinical Trials
- Phase IIIb/IV: Peri/Post-Approval
- Registries
- Observational Research
- Expanded Access Programs
- Safety Studies
- Clinical Logistics

PAREXEL CONSULTING

- Integrated Product Development
- Regulatory Outsourcing Services
- HERON™ Commercialization
- Medical Communications
- Strategic Compliance and Risk Management



LEARN MORE AT PAREXEL.com

WHEREVER YOUR JOURNEY TAKES YOU, WE'RE CLOSE BY.

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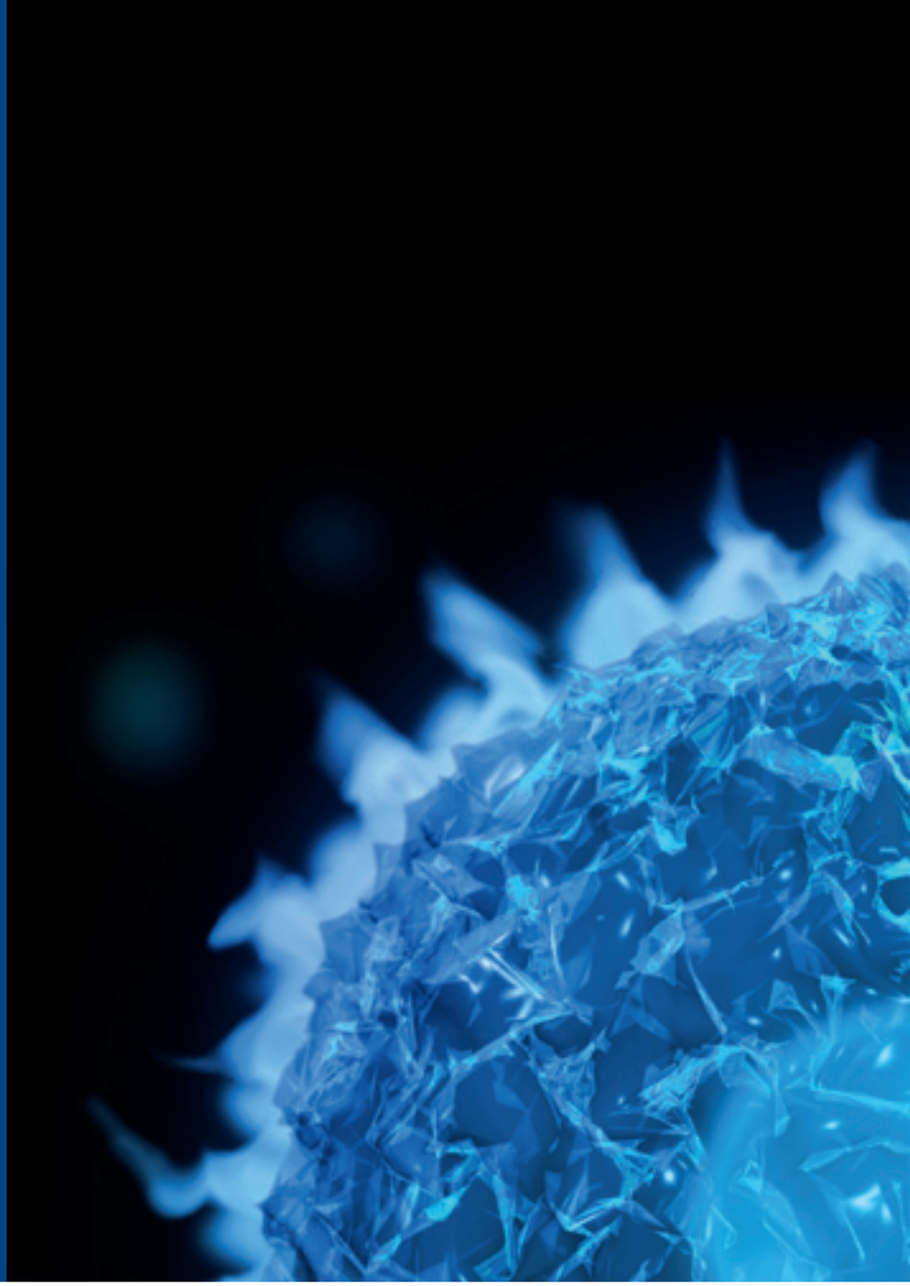
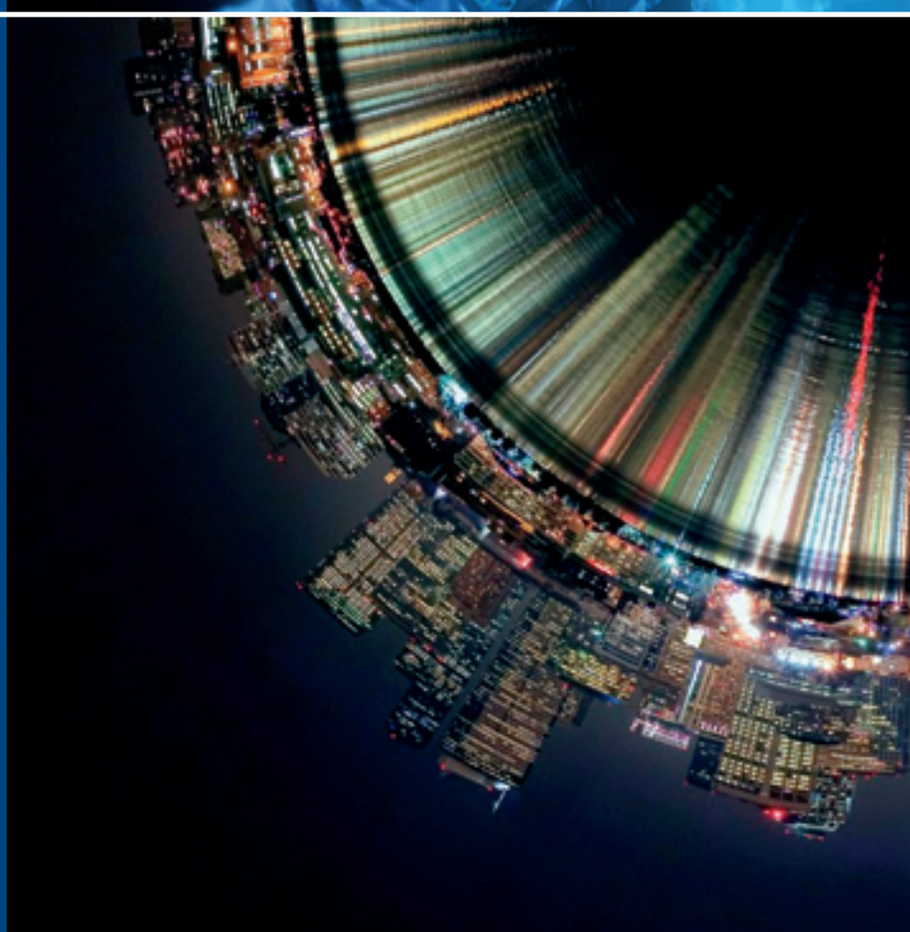
Our regional leads are always available for a conversation.





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Cover & back cover: organic cell (top image), Seattle cityscape (bottom image)