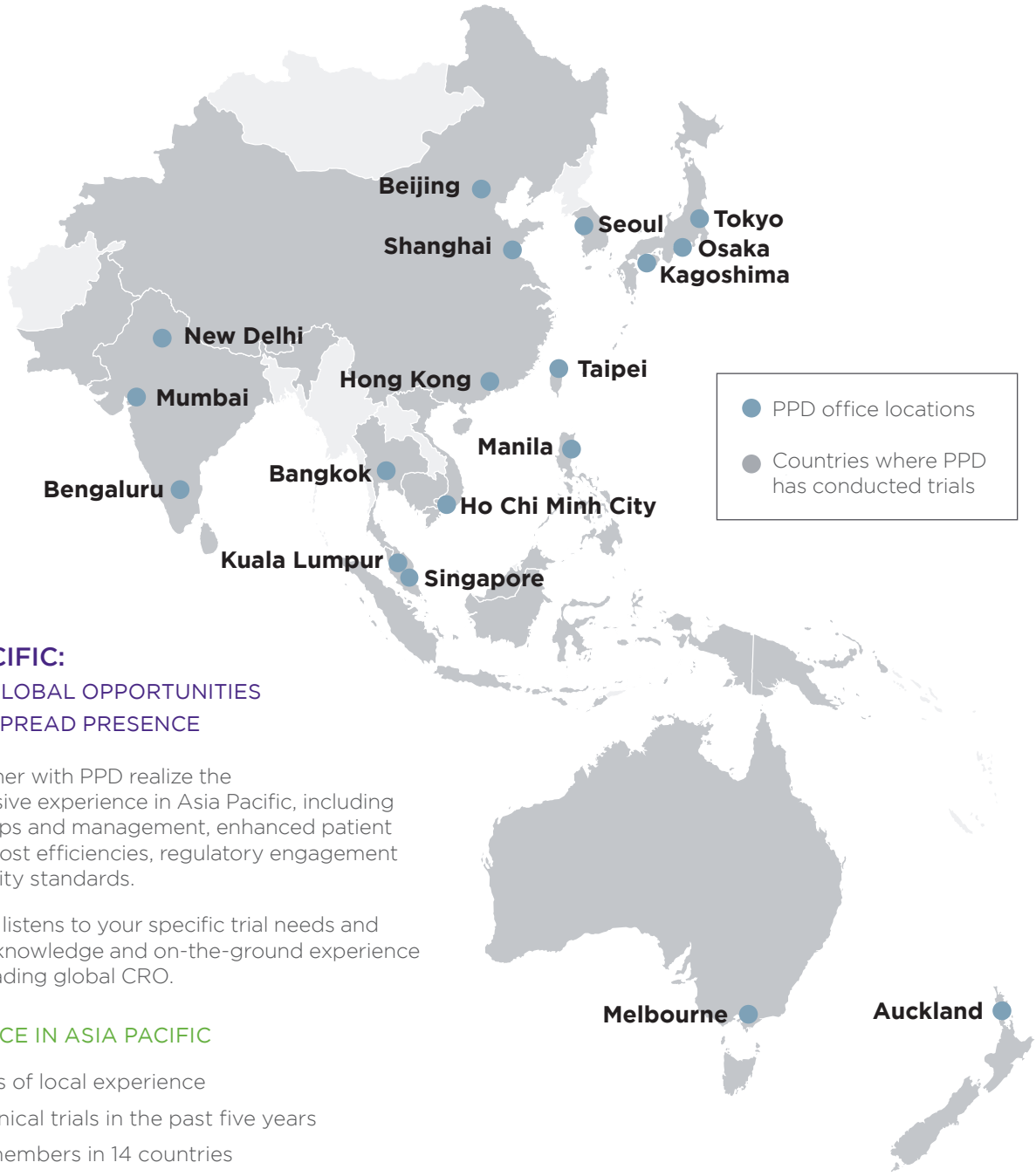
PPD®

LOCAL EXPERTISE

ASIA PACIFIC

SEIZING OPPORTUNITIES THROUGH IN-DEPTH LOCAL KNOWLEDGE

With more than 15 years of extensive local experience in Asia Pacific and global integration of operational functions, PPD is the partner of choice in the region.



PPD IN ASIA PACIFIC:

CAPITALIZING ON GLOBAL OPPORTUNITIES REQUIRES A WIDESPREAD PRESENCE

Companies that partner with PPD realize the benefits of our extensive experience in Asia Pacific, including strong site relationships and management, enhanced patient selection, increased cost efficiencies, regulatory engagement and outstanding quality standards.

PPD is a partner who listens to your specific trial needs and provides the quality, knowledge and on-the-ground experience you expect from a leading global CRO.

THE PPD DIFFERENCE IN ASIA PACIFIC

+ More than 15 years of local experience
+ More than 600 clinical trials in the past five years
+ 1,700 PPD team members in 14 countries
+ Complete range of globally integrated services
+ Clinical joint venture and large team in Japan

A COMPREHENSIVE RANGE OF SERVICES FOCUSED ON YOUR SUCCESS

+ Project management
+ Clinical management
+ Regulatory affairs
+ Pharmacovigilance
+ Biostatistics
+ Data management
+ Late stage research
+ Central labs

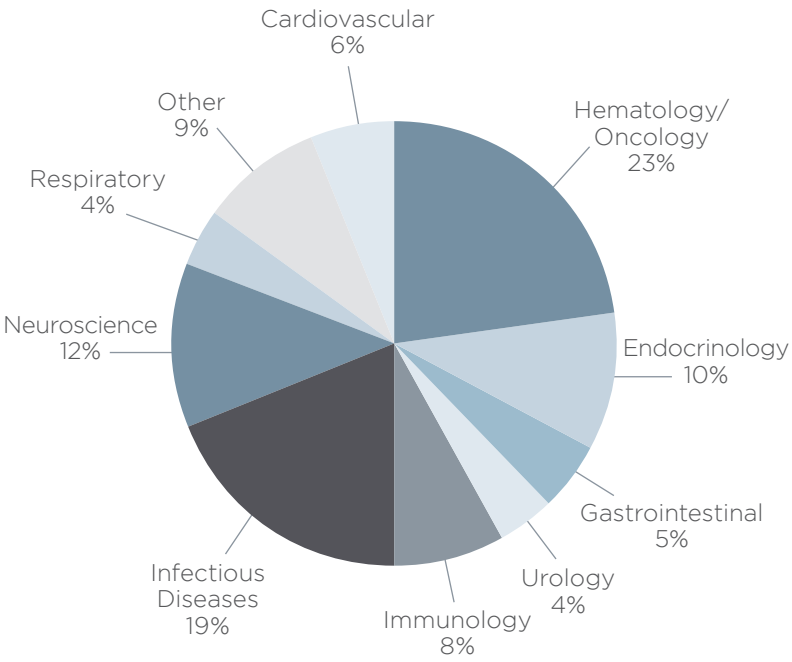
COMPREHENSIVE THERAPEUTIC EXPERTISE. **EFFECTIVE SITE MANAGEMENT.**

At PPD, we are committed to ensuring your clinical trials in Asia Pacific are successful from beginning to end. We proactively manage your sites to ensure the highest standards of quality and safety.

When it comes to site selection and management, PPD engages an established and extensive network of resources in Asia Pacific. We are passionate advocates of your study, constantly promoting your trials to investigator teams that have experience in a wide array of therapeutic areas. We maintain strong and positive relationships with our investigators and their site staff to ensure efficient enrollment and high-quality conduct of your trials.

As with all PPD clinical development operations around the world, our engagement with Asia Pacific-based sites is completely transparent, globally aligned and adheres to the strictest clinical quality assurance standards and standard operating procedures (SOPs). You'll receive prompt and detailed insight regarding any emerging risks or concerns regarding your trial—providing you with ample time to make any necessary adjustments.

EXTENSIVE CLINICAL EXPERIENCE IN ASIA PACIFIC



NAVIGATING MULTIPLE REGULATORY STRUCTURES WITH A SINGULAR COMMITMENT TO EXCELLENCE

Developing an effective strategy to address the complexities of the Asia Pacific regulatory environment is certainly a challenge. But, with PPD's expertise, it's one that is completely surmountable.

PPD has one of the most experienced regulatory affairs teams in Asia Pacific, and we deliver many benefits to our clients, including:

+ More than 40 regulatory affairs staff members located throughout the region
+ Strong local relationships
+ In-depth knowledge of strategic and tactical regulatory pathways
+ Frequent training to ensure consistent quality
+ Alignment with PPD's global regulatory team

PPD has proven expertise in creating and deploying a comprehensive development plan and regulatory strategy for clients that includes advanced planning, local knowledge and strong professional relationships with regulatory agencies.

EXPERT RESOURCES FOR CLINICAL STUDIES IN JAPAN

PPD has joined up with SNBL to create a new clinical joint venture in Japan, PPD-SNBL. We are one of the largest clinical outsourcing CROs in Japan. With strong knowledge of the Japanese clinical trial landscape, you can rely on PPD-SNBL to deliver efficient, high-quality clinical trial outcomes whether you are conducting Japan-only studies or whether Japan is a component of a larger global study.

+ Full-service Phase I-IV clinical trial capabilities
+ Nearly 20 years of clinical trial experience in Japan
+ 400 staff in offices in three cities in Japan
+ More than 150 clinical trials conducted in the past five years
+ Local knowledge and experience of a strong Japanese company established more than 50 years ago coupled with the geographical reach and top-tier technology and processes of a leading global CRO.

LABORATORY SERVICES IN ASIA PACIFIC

CENTRAL LABS

Timely access to clean, accurate data is essential to the success of any clinical trial. PPD® Laboratories ensures you'll have that no matter where in Asia Pacific your trial is located.

PPD Laboratories provides a complete range of integrated clinical trial management and laboratory services throughout the Asia Pacific region. You'll benefit from increased efficiencies and greater study flexibility through:

+ Strategically located laboratories that ensure uninterrupted coverage to all regions
    - Highland Heights, Kentucky
    - Brussels, Belgium
    - Shanghai, China
    - Singapore
+ PPD Clicks™ technology—secure website to view and analyze the most current lab results
+ Integrated CRO and central lab data through a proprietary, automated platform, Preclarus™

CONSISTENT RESULTS ACROSS THE GLOBE

+ PPD Laboratories has a single, global database which eliminates the need to integrate and harmonize data
+ Our laboratory methodology, instrumentation, reagents and processes are standardized across all of our central lab locations
+ Assays are extensively cross-validated by all central lab locations to produce combinable data

For more information, please contact us
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