

. Our Services

We support international research and development in Asia

In 1999 we became the first Japanese CRO to establish a subsidiary in China and have since started planning, regulatory submissions and project management of clinical trials in Asia. We have numerous medical professionals and expert staff well versed in clinical trial regulations who provide high quality clinical trial services in China and Southeast Asia. We currently support local submissions and clinical investigations for submission in Japan and clinical research for EBM through partnerships with EPS China Co., Ltd. (Shanghai, Beijing, Guangzhou), EPS Singapore (Ever Progressing System Pte Ltd.), and other partners.

Through partnership with leading world CROs based in Europe and the U.S., we also support global development of domestic and overseas pharmaceutical companies. Based on our clinical development know-how and experience cultivated as a CRO, in the future we intend to reinforce such consultation capabilities as licensing and marketing strategies, with a view toward business expansion to pharmaceutical companies and bio-venture companies in Asia, especially new businesses in product development utilizing Chinese market characteristics, in order to further enrich our services.

Major Services

- Regulatory consultation/development of clinical trials
- Support for development of documents for regulatory submission and regulatory application on behalf of clientele
- Development of trial-related documents such as protocol and CRFs
- Drug randomization
- Monitoring
- Data management
- Statistical analysis
- Medical writing
- Various marketing surveys and marketing promotion consultation

EPS overseas network



Countries and regions to which we can respond:

China, Hong Kong, Taiwan, Korea, Singapore, Malaysia, US, South Africa

ADVANTAGE of EPS

- 1 The number of staff totals over 100 employees with medical and pharmaceutical career backgrounds and experience in DM and statistical analysis.
- 2 Our company is one of the largest CROs in China and, with its unique web-based case enrollment and data management systems, is capable of large-scale clinical trials intended for EBM.
- 3 Local staff can speak both Japanese and English. They can give detailed responses appropriate to client requests.
- 4 Utilizing our extensive network with local administration and medical sites, we realize efficient clinical trials.

Our contract experience includes achievements in:

[Therapeutic areas]

Cardiovascular drugs, ophthalmic drugs, metabolic drugs, CNS drugs, PNS drugs, allergic drugs, external dermatological drugs, tumor drugs, antibiotics, chemotherapies (antivirus agents), digestive organ drugs, revitalizers, etc.

[Services]