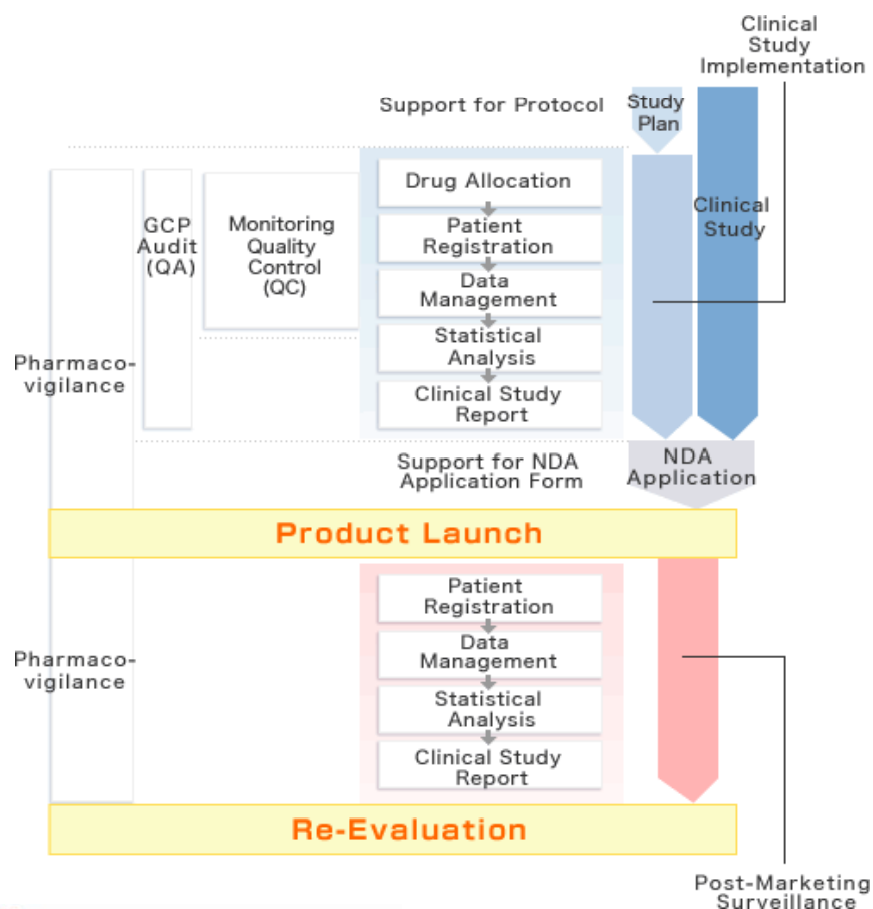


New Drug Development Services

We Provide Speedy and High Quality Services for New Drug Development

EPS supports all services-related data management, statistical analysis, monitoring, protocol designs, regulatory application and post-marketing surveillance in clinical studies. We are also utilizing the capabilities of EPS to support Asian and Global-studies with high quality standards and providing clinical development solutions in a timely manner.



Supported-Studies

- Clinical Study
- Post-Marketing Surveillance
- Investigator-Initiated Study
- Extensive Epidemiology Surveillance
- Clinical Research

Supported-Services

- Monitoring
- Patient Registration Center Service
- Data Management
- Statistical Analysis
- Medical Writing
- Pharmacovigilance
- GCP Audit
- Consultation, Regulatory and Others
- System-related Clinical Development and ASP Service

Supported-Areas

- Japanese-Study
- Asian and Global-Study

Enhanced Solutions

● Global Clinical Development Services

We provide Asian and Global studies with high quality standards through alliances with leading global CROs world wide and EPS networks in Asian countries.

● Data Center Services

In extensive post-marketing surveillance, we provide client-specific data management services with EPS know-how and human resources.

● Offshore Data Management Services

Data management is operated at our subsidiary company in China with the same high quality and speed guarantee as in Japan.

ADVANTAGE of EPS

- Full-services for product development from pre-clinical studies to post-marketing surveillance.
- Support mega-study with enriched experiences in oncology and cardiovascular diseases.
- Having specialists with expert knowledge and standardized procedures, we guarantee high quality standard services.
- Through EPS' overseas network, we support Asian and global studies.
- With experienced know-how and comprehensive systems we are able to develop clinical study plans.