Your Trusted Service Partner for the U.S., Japan, and Asia

Better Medicine Sooner

The First and Largest CRO in Japan Offers You Services Globally

CMIC was founded in 1992 by Dr. Kazuo Nakamura as the first contract research organization (CRO) in Japan.

To enable pharmaceutical companies to develop better medicine sooner, we expanded our solutions to include contract development and manufacturing (CDMO), contract sales (CSO), and healthcare services and established an innovative pharma model (IPM) to bring highly desired treatments to the Japanese market. We strive to be a Pharmaceutical Value Creator (PVC), spanning our services across the entire drug development value chain, and meeting global customers' needs in the U.S., Japan, and broader Asia.

Largest CRO in **JAPAN** with 1,200+ CRAs

Involved in nearly 80% of new drug development in **JAPAN**

Work with Top 10 global pharma companies

Connected with 2,600+ clinical sites in **ASIA-PACIFIC**

The First CRO in **SOUTH KOREA** with experience in 300+ Studies

Experienced in 4,500+ Protocols

Among Top 3 largest CDMOs in **JAPAN**

The Only CRO in **JAPAN** with CMC, bioanalysis, and nonclinical services

2nd Largest CSO in **JAPAN** with over 650 contract sales reps and MSLs

Largest regulatory consulting and medical writing team in **JAPAN** with 120+ Experts

Completed 300+ oligonucleotide assays in the **U.S.**

12+ Years of continuous commercial manufacturing and drug development services in the **U.S.**

Our Tailored Solutions

CMIC offers tailored solutions for pharmaceutical companies, medical device manufacturers, academia, bio-ventures, and medical institutions. Using our collective strengths, we offer a broad range of services from pre-clinical research to practical applications.

Japanese Market Entry

Providing full pharmaceutical capabilities for overseas business partners to enter the Japanese market.

- In-country clinical caretaker (ICCC)
- Marketing authorization holder (MAH) services

Asia Clinical Trials

Supporting our clients' clinical trials in 13 countries in the Asia-Pacific region.

- Consulting services
- Regulatory affairs
- Project management
- Clinical operations

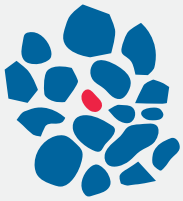
Medical Device & In Vitro Diagnostics

- QMS organization setup
- Design and development
- Nonclinical studies
- Clinical trials
- Approval/certification support
- Reimbursement support
- Post-marketing surveillance

Oncology Drug Development

Highly specialized oncology experts support your drug development.

- Consulting services
  - Strategic development planning
  - Study design optimization
- Regulatory affairs
- Clinical trials
- Site and patient support
- CSO services



Orphan Drugs

- Regulatory affairs
- Manufacturing, packaging/labeling
- Quality management
- Safety management
- Medical affairs
- Marketing
- Supply chain management
- Sales and distribution

Regenerative Medicine

Established systems ready to implement and form development strategies and support every stage of the regenerative medicine life cycle.

- Consulting services
- Regulatory affairs
- Nonclinical studies
- Clinical trials

The Only End-to-End Solution Partner in Japan

Pre-Clinical

Clinical

Consulting

- Drug development consulting
- Japan/foreign market entry
- Medical devices consulting
- Out-licensing to other pharma

Regulatory Affairs

- Regulatory submissions and correspondence

Laboratory

- Chemistry manufacturing and control (CMC) and GMP analytical services
- GLP bioanalysis and analytical services
- Nonclinical studies (toxicity, safety, and efficacy)

Formulation Design

- Formulation, process, and analytical development

Clinical Trials

- ICCC/local country agent
- Project management
- Clinical operations
- Data management
- Pharmacovigilance
- Quality assurance and control
- Statistical analysis
- Medical writing
- Patient recruitment and retention

Investigational Product Manufacturing

- Formulation, process, and analytical development
- Early-to-late-phase manufacturing
- Packaging and labeling
- Release and stability tests
- Technology transfer
- Antibody active pharmaceutical ingredients manufacturing

Laboratory

- CMC and GMP analytical services
- GLP bioanalysis and analytical services

Site and Patient Support

- Patient call center and help desk
- Clinical research coordinator (CRC) services
- Clinical site administration support

Medical Affairs

- Evidence generation
- QOL study
- Medical affairs consulting
- Medical science liaison (MSL) talent solutions
- MSL training and assessment
- Patient support program

Other Services and Products

- Orphan drugs

As the only Pharmaceutical Value Creator (PVC), CMIC Group can take your product from pre-clinical, to clinical, to filing/approval and through post-marketing studies seamlessly.

Whether in the U.S., Japan, or elsewhere in Asia, we are your dedicated, customer-centric, full-service provider across the full drug development continuum.

Filing/Approval

Post-Approval

- In vitro diagnostics consulting
- CMC Consulting
- Regenerative medicine consulting
- Product life cycle management

Consulting
- MAH related services
- National health insurance (NHI) pricing
- Health technology assessment (HTA) related consulting

Medical Writing

Commercial Supply
- Commercial manufacturing
- Packaging and labeling
- Release and stability tests
- Antibody active pharmaceutical ingredients manufacturing

Health Economics and Outcomes Research

Medical Affairs
- Evidence generation
- Medical affairs consulting
- Medical science liaison (MSL) talent solutions
- MSL training and assessment
- Materials review

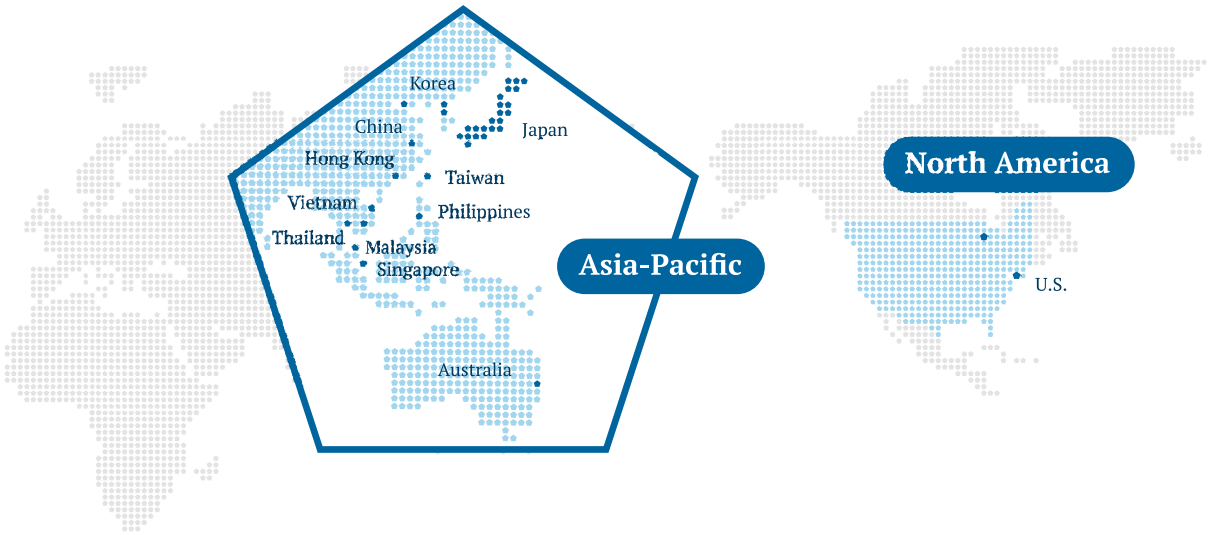
Patient and Healthcare Services
- Call center
- Patient support program
- Digital healthcare channel service: harmo
- Prescription drug database services
- Healthcare website: HelC+
- Self-test kit: SelCheck

Post-Marketing Surveillance and Clinical Research

Sales and Marketing
- Sales/medical representative solutions
- Remote detailing services
- Medical device sales support
- Customer service representatives
- Clinical/nurse educator

Agile Access to the World's Major Pharmaceutical Markets

A variety of services can be provided, worldwide, with CMIC facilities or modern technology.



Services		U.S.	Japan	Korea	Singapore	China	Taiwan	Malaysia	Hong Kong	Philippines	Vietnam	Thailand	Indonesia	Australia	New Zealand
Pre-Clinical	Consulting	⬟	⬟												
	Laboratory	⬟	⬟												
	Formulation Design	⬟	⬟												
Clinical	Clinical Trials		⬟	⬟	⬟	⬟	⬟	⬟	⬟	⬟	⬟	⬟	⬟	⬟	⬟
	Investigational Product Manufacturing	⬟	⬟	⬟											
	Laboratory	⬟	⬟												
	Medical Affairs		⬟												
	Regulatory Affairs		⬟	⬟	⬟	⬟	⬟	⬟	⬟	⬟	⬟	⬟	⬟	⬟	⬟
	Site and Patient Support		⬟												
Filing/Approval	Consulting		⬟												
Post-Approval	Commercial Supply	⬟	⬟	⬟											
	Medical Affairs		⬟												
	Patient and Healthcare Services		⬟												
	Post-Marketing Surveillance and Clinical Research		⬟	⬟	⬟	⬟	⬟	⬟	⬟	⬟	⬟	⬟	⬟	⬟	⬟
	Sales and Marketing		⬟												

*Full range of services for development stage may not be available for some locations.

CMIC's Mission

CMIC is an innovative and unique provider of high-quality solutions for the healthcare industry. We create value by accelerating the access to therapies that improve patients' lives.

CMIC's Vision

To advance the innovation of products and solutions that will empower people worldwide to achieve greater health and wellbeing.

CMIC's Values

W & 3C

WELLBEING: Fully Live Every Moment
Challenge: Liberate opportunity by changing our vantage point
Change: Transform without seeking refuge in conventional wisdom
Communication: Proactively reach out to people and society

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