



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 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), site support, and market solutions, supporting global companies to bring highly desired treatments to the Japanese market.

Leveraging our vast experience and expertise in the medical industry, CMIC also offer solutions using a new ecosystem of healthcare to support individuals and local governments.

We strive to be a **Personal Health Value Creator (PHVC)** to meet our global customers' needs in the U.S., Japan, and broader Asia.

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

One of the
Largest
CROs
in **JAPAN** with
leading
ICCC studies

Among
Top 3
largest CDMOs
in **JAPAN**

Connected with
4,000+
clinical sites in
ASIA-PACIFIC

The Only
end-to-end
JAPAN
MARKET ENTRY
solution
provider

Involved in nearly
70%
of new
ONCOLOGY
drug development
in **JAPAN**

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

Completed
400+
oligonucleotide
assays in
the **U.S.**

Largest regulatory
consulting and
medical writing
team in **JAPAN** with
150+
Experts

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

The First CRO in
SOUTH KOREA
with experience in
600+
Studies

2nd Largest
CSO in **JAPAN**
with over
750+
sales reps,
nurses, and MSLs



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

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

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
- Local government solutions



CMIC Group can take your product from pre-clinical, to clinical, to commercialization 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

- In vitro diagnostics consulting
- CMC Consulting

Post-Approval

- Regenerative medicine consulting
- Product life cycle management

Consulting

- MAH related services
- National health insurance (NHI) pricing

Medical Writing

Commercial Supply

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

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
- Healthcare communication channel: harmo
- Prescription drug database services
- Healthcare website: HeIC
- Self-test services: SelCheck
- Specialty care support platform: nanacara

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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