

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

Better Medicine Sooner



Leading CRO in Asia-Pacific Offering Broader Services Globally

Founded in 1992, CMIC pioneered Japan's first contract research organization (CRO) and later became the country's first site management organization (SMO). We also extended our footprint across Asia as the first CRO to operate in South Korea.

To enable pharmaceutical companies to develop better medicine sooner, we expanded our solutions to include contract development and manufacturing (CDMO), consulting, 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.



Our Specialized Solutions

CMIC offers 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.

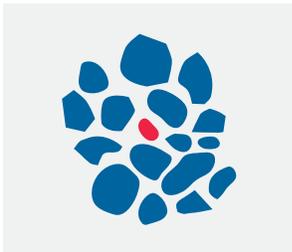
- Consulting services
- Regulatory affairs
- Project management
- Clinical operations



Oncology Drug Development

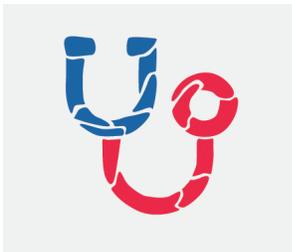
Highly specialized oncology experts support your drug development.

- 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



Medical Device & In Vitro Diagnostics

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



Cell & Gene Therapy

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 Pharmaceutical Solution Partner in Japan

Pre-Clinical

Clinical

Consulting

- Drug development consulting
- Japan/foreign market entry
- Interview/research with physicians, paramedical, patients and families
- 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

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

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



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.



- In vitro diagnostics consulting
- CMC Consulting

- 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

Site and Patient Support

- Call center
- Patient support program

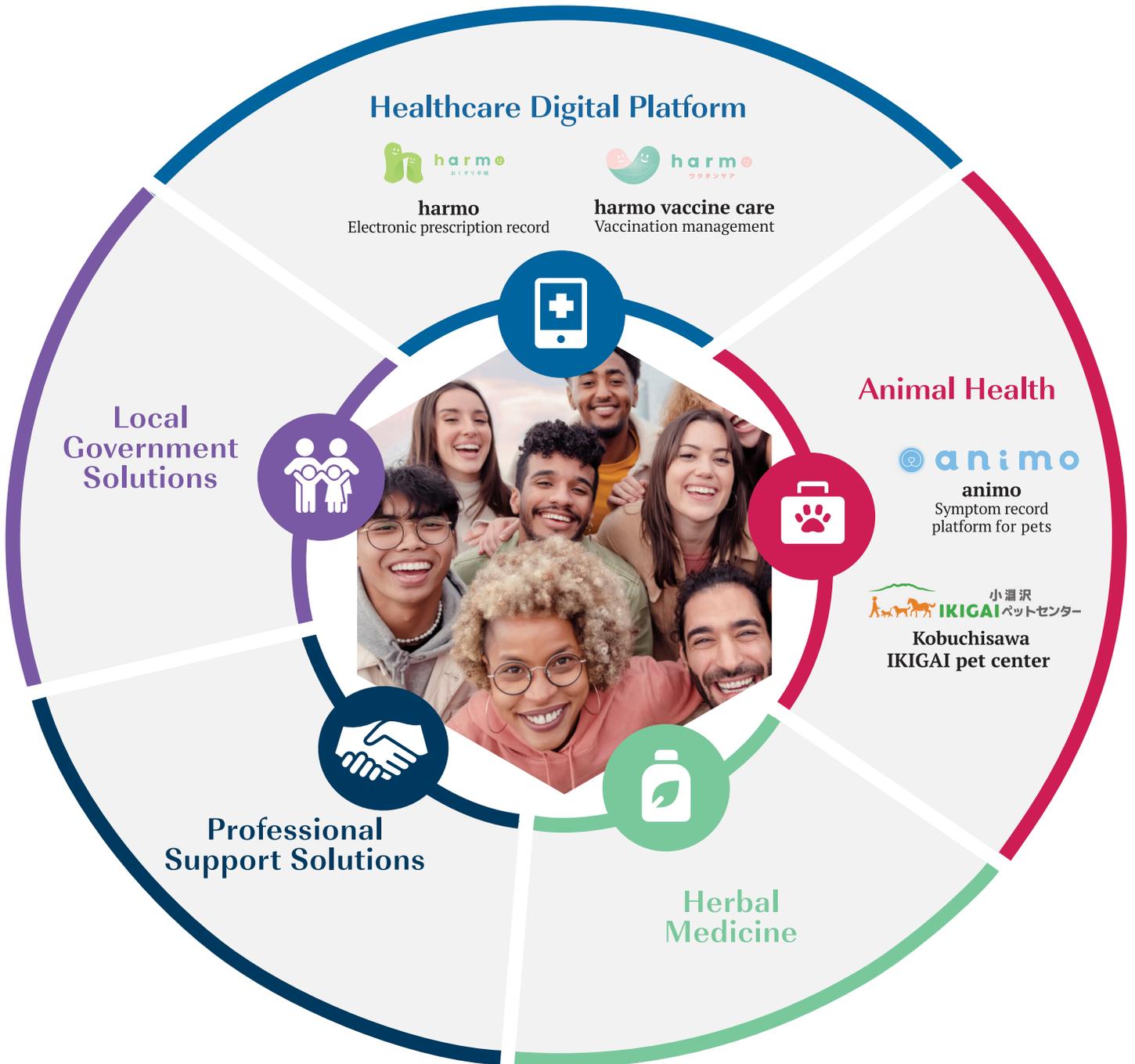
Post-Marketing Surveillance and Clinical Research

Sales and Marketing

- Sales/medical representative solutions
- Remote detailing services
- Sales support
- Customer service representatives
- Clinical/nurse educator

Innovative Healthcare Solutions

CMIC is committed to providing patient-centered healthcare solutions that meet the evolving needs of various diseases through advanced technology, including digital/IoT and contact center services.



Agile Access to the World's Major Pharmaceutical Markets

CMIC supports clinical trials in Asia-Pacific, Japan market entry, and bridging non-clinical and manufacturing needs between the U.S. and Japan.

- Japan
- Korea
- China
- Hong Kong
- Taiwan
- Singapore
- Malaysia
- Philippines
- Thailand
- Vietnam
- Indonesia
- Australia
- New Zealand



North America

- New Jersey
- Illinois





CMIC Group

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