



Contract Research Organization, Your End-to-End Solutions Provider [Q](#) [☰](#)

Contract
Research
Organization



Contract
Development &
Manufacturing
Organization



Healthcare



Contract Sales
Organization



Japan Market
Entry



Visit Our Virtual Booth

Speak with an Expert

Pharmaceutical Development Services

CMIC Group pioneered the Contract Research Organization (CRO) business in Japan more than 30 years ago. Since then CMIC has continued as a leading innovator in the industry and has expanded services to encompass the entire pharmaceutical value-chain, including: contract development and manufacturing, healthcare solutions, sales and marketing, Japan market entry solutions and more. We pride ourselves on meeting customers where they are and with exactly what they need to move their products forward.

View our services below by selecting the phase of drug development that you are working in:



Pre-Clinical



Clinical
Phase I-III



Approval



Post-Approval, Sales &
Marketing



[Consulting Services](#) >

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Agreement Targeting New
Value Creation in Medical and
Healthcare Field

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Japan's first and largest CRO
continues 30-year evolution to
help global sponsors navigate
the complex local market

[Access Full Interview](#)

Bioanalytical Method
Validation: Challenges and
Opportunities of a Globally
Harmonized System

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Our Tailored Pharmaceutical Solutions

CMIC, Japan's first and largest Contract Research Organization, 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 preclinical research to practical applications.



Japanese Market Entry

Providing full pharmaceutical capabilities for overseas business partners to enter into Japan market.

[Learn More](#)



Asia Clinical Trials

Supporting our client's clinical trials throughout the Asia Pacific region. Providing solutions for a range of therapeutic areas, registry study, observational study, phases II/III clinical trials, phase IV clinical trials, post-marketing surveillance.

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Medical Device & IVD

Consultation services on regulatory issues for the latest medical devices and in vitro diagnostics (IVD).

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Oncology Drug Development

Specialized oncology staff assists with strategic development planning, including regulatory interaction, conducting clinical studies and post-marketing promotion for companies that wish to enter into the field of oncology.

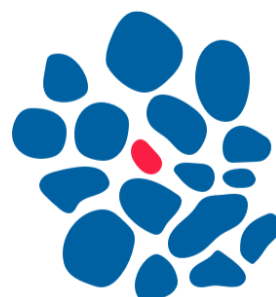
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Cell & Gene Therapy

Established systems ready to implement and form development strategies and support every stage of the regenerative medicine lifecycle, including regulatory consulting, various nonclinical and clinical studies, and post-marketing.

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Orphan Drug

OrphanPacific is a pharmaceutical company within CMIC Group, which provides wealth of experience in the field of orphan drugs: regulatory affairs, supply chain management, quality management and marketing strategy planning.

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CMIC Contract Research Organization: Latest Resources



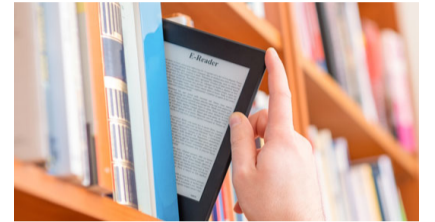
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