We offer professional, integrated services for new product development from A to Z service

Regulatory Consulting | Clinical Trials | Global Market Access



A to Z service

From R&D to Clinical trial, licensing, commercialization stages and late phase study, we provide full scope of one-stop solution and integrated consulting.



Experience in projects

We have performed more than 2,200 projects management in clinical trials since 2000 (Including global sponsor's projects) More than 1500 IND & NDA projects in Drug and medical device (including US FDA services)



Global network

DreamCIS / MediTip family has a big global network via our another family, Tigermed group. Based on our global capability, we can provide the service that our customers want.



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About Us

Core expertise

2200+

Clinical Trial Project Management

1500+

IND & NDA services (Drug & medical device)

50+ RA

Biggest regulatory Team

51 Countries

Global network



We, as a family, provide innovative A to Z solutions as your global partner.



Founded in 2000, DreamCIS, with 2000+ project experience, provides full-scope services including Phase 1-4, rPMS to NIS. After DreamCIS was acquired by Tigermed in 2015, DreamCIS enhanced its global network and became a step closer to global CRO by SOP integration with Tigermed. DreamCIS has reinforced its presence as Korean TOP clinical CRO by listing on KOSDAQ market as the industry's first in 2020. In 2022, DreamCIS welcomed MediTip, No.1 IND/NDA consulting service provider in Korea, as a family to provide A to Z, one-stop service to customers, to enlarge its business area and to continue its growth as top global CRO.



Founded in 2007, Meditip guides you into Korean and global healthcare market with intelligence, in-depth knowledge and experiences. Meditip has rich experiences in successfully collaborating with global leading bio-technology suppliers for Korea and global market entry issues, pre-market regulatory affairs, post-market compliance affairs, etc. As Korean TOP IND/NDA consulting service provider, Meditip is ready to provide high quliaty of intelligence that is crucial to your business success in Korean and global market.



LCS, founded in 2017, is a non-clinical consulting and project management agency. LCS offers the best service collaborating with professional research organizations. We provide R&D service consulting for lead optimization, non-clinical consulting for IND/NDA of new drug and biomedicine, etc. LCS promises to lead our clients' successful business through our various experiences, know-how, professional resources and special network with cooperative organizations.

One Dream. One Family.



Our A to Z Service



Pre-Clinical

- Strategy consulting for nonclinical development
- · Regulatory consulting of formulation/CMC
- · Review pre-clinical study protocol and report
- Review and analysis between nonclinical and clinical

Clinical Trial I-III

- · IND/IDE submission
- Full Scope of Clinical Development from Phase 1 to Phase 3
- · Full scope Medical Device Clinical Development
- · Clinical Strategy Development Plan
- · Global Multi-national Trials
- · Pharmacovigilance & Medical Writing
- · Data Management & Statistics

Registration

- · NDA/BLA application
- · CTD writing and NDA/BLA filing
- · RMP/PMS development
- Feasibility assessment
- Strategy development for approval of product registration

Market access & ICC

- · PE study, Reimbursement
- Market forecast and license out
 ICC(In-Country Caretaker) service
- · Support SOP for GIP

Late Phase Study

- Full scope of Phase 4 Clinical Trials
- Post Marketing Surveillance
- Non-intervention studies & Observational studies / safety surveillance
- Long term follow up observational Study
- RMP Service & Pharmacovigilance
- Real World Evidence
- Registry Study
- HIRA Data Analysis
- Retrospective Chart Review

Global Access Service —

- \cdot Oversea's Clinical Trials in
- EU. US. China etc.
- Drug development
- Medical Device

Our Global Coverage



