- Maximize Commercial Potential of Product with Global Strategy
- Faster Time to Market, Maximize Commercial Opportunity
- Achieve Global Compliance, Reduced Risk
- Stay in Control with Centralized Management
- 100% Approval Success

- Clinical and Regulatory Experts with decades of experience with FDA, EMA, ANVISA, CFDA, PMDA, Cofepris
- Regional Offices in USA, Europe, Brazil, China, Japan, Mexico
- Clinical Trial facility in Brazil

## iCRMAX

*Integrated Clinical, Regulatory, Market Access Xcellence Platform*

Learn More

- *Acceptance of Foreign Clinical Trial Data per 21 CFR part 312 IND regulations and 21 CFR part 812 IDE regulations.] Under 21 CFR 312.120(c)(1)*
- *Acceptance of Foreign Clinical Trial Data by NMPA/CFDA, Technical Guiding Principles for the Acceptance of the Overseas Clinical Trial Data of Drugs, July 2018*