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BVMA member since 1996
Audits passed in 2000, 2003,
2006, 2009, 2012 and 2015

HISTORY AND MISSION

For more than two decades SynteractHCR has provided support to clients in all phases of clinical development, across multiple therapeutic areas. Our standards of integrity and reliability have been mainstays throughout our history, as proven by our established track record of repeat and referral clients. Our mission is to take time and cost out of the drug development process so we can get successful therapies to the patient population that needs them sooner.

HIGHLIGHTS

SynteractHCR is a full-service CRO with a two-decade track record supporting biotechnology, medical device and pharmaceutical companies in all phases of clinical development. We have conducted Phase I-IV studies on six continents and in more than 50 countries. Our expertise in multiple therapeutic areas includes notable depth in oncology, CNS, infectious disease, endocrinology and other indications. Intelligent Clinical Development (ICD+) is the approach we use to leverage our deep clinical and therapeutic expertise, appropriate technology, and optimized process execution to drive solution-oriented efficiencies on a global scale. ICD+ allows sponsors to get to decision points faster, while providing high quality, consistent standards and clear results internationally.

SERVICES

»Shared Work – Shared Vision« is the philosophy behind the way we do business. It's a promised standard underlying our full suite of CRO services that includes expertise, mentoring, custom planning and collaborative support.

- **Project Management:** Our methodology incorporates standardized processes and metrics for reviewing project progression, risk assessment-mitigation and completion.
- **Clinical Operations:** Experienced clinical managers and clinical research associates oversee each study's clinical processes to ensure proper planning, conduct, patient safety, and data quality, while fostering good communication between study sites and sponsor.
- **Data Management:** Excellent planning and organization allow us to meet aggressive timelines in all phases of clinical development with accuracy levels consistently above 99.9%
- **IVRS/IWRS:** We offer full-featured global Interactive Voice and Web Response Systems for your clinical trial operations, through our validated 21 CRF Part 11 compliant proprietary system.
- **Medical & Regulatory Affairs:** Our services span the full range of product development, including training, performed by senior regulatory professionals with international experience in pharmaceutical, diagnostic, device and biotechnology industries.
- **Medical Writing:** Whether it's a first clinical protocol, a pivotal trial protocol or associated documents such as the ICF, the investigator's brochure, or a full IND/IDE, our team brings the therapeutic, clinical, regulatory, and writing expertise needed to ensure the success of your clinical trial from beginning to end, design phase to final regulatory submission.