CENTAUR CLINICAL

Tests & Trials for your CE marking

Preclinical Testing

Centaur Clinical performs preclinical testing of your devices in a modern well-equipped testing laboratory in accordance with ISO10993 and Good Laboratory Practices.

Clinical Trials

Specialized in Clinical Operations, Centaur clinical sets up clinical trials, from study design to post market phases, in accordance with Good Clinical Practices.

Medical Writing

Centaur Clinical writes up the regulatory documents necessary for your CE marking following the MDR 2017/745 and the EU MEDDEV 2.7 rev.4.

A multilingual team with a dedicated and optimized support

Centaur Clinical helps medical device and diagnostic manufacturers in their CE marking process, from conducting preclinical testing to clinical trials and post-market studies.

Thanks to our multilingual team and our international network of partners, Centaur Clinical is able to set up rapidly optimized studies in specific areas such as Cardiology, Oncology and Intensive Care.

Since 2019, Centaur Clinical has been engaged in the ISO 13485 certification process, and services are already provided in accordance with ISO 14155 and GCP.

Tél. 09 67 08 56 76 • info@centclin.com • www.centaurclinical.com

Pôle d'activités Puits Morandat - 1480 avenue d'Arménie - 13120 Gardanne

Centaur Clinical Services for your CE marking

CENTAUR CLINICAL

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Preclinical testing

- Cytotoxicity
- Hemo-compatibility
- Genotoxicity
- Pyrogenicity
- Immuno-genicity
- **Technical file drawing up**

Clinical Trials

- Study Design
- Protocol Writing
- Site Selection
- Feasibility study
- Regulatory Submissions
- Implementation
- Study follow up
- Data Analysis
- **Clinical Study Report**

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Study Preparation | Data Management | Monitoring | Regulatory files