

YOUR CLINICAL RESEARCH SOLUTION

A BOUTIQUE SITE MANAGEMENT ORGANIZATION



YOUR PARTNER IN CLINICAL RESEARCH

SciMedBE was founded to answer the needs of the pharma and medical devices industry, looking for a flexible Site Management Organization (SMO) with therapeutic excellence in Anesthesiology, Critical Care, Pain and Perioperative Medicine.

Headquartered in Belgium, the heart of Europe, SciMedBE capitalizes on Belgium's streamlined regulatory environment, trained professionals, compliant patients and motivated clinical sites. As a Site Management Organization we assure the fastest possible start-up of our research sites within our network. In 2017, SciMedBE was the top enrolling network out of 20 sites in FDA related trials.





ADVANTAGES OF A DEDICATED SMO

REDUCTION OF TIME & COSTS OF DRUG DEVELOPMENT BY

- ✓ Clinical Research Network in Belgium and Europe (Denmark, Croatia, Slovenia)
- ✓ Standardized Service Agreements for each customer
- ✓ Improved communication between multiple investigational sites and the client
- ✓ Reduction in administrative issues
- ✓ Faster study start-up
- ✓ Providing on-site support
- ✓ Enhancing patient enrollment: rapid recruitment of qualified subjects
- ✓ Direct contact with decision makers and management



OUR TEAM



Ingrid Meex, MSc, PhD
Research Coordinator



Catherine Vandepitte, MD
Research Consultant



Prof. Admir Hadzic, MD, PhD
President



Sam van Boxstael, MD
Research Consultant



Marijke Cipers, Bsc
Clinical Research Nurse



Gülhan Özyürek, MSc
Clinical Research Associate



Sigrid Christiaens, MSc
Clinical Research Associate



Tamara Sals, MSc
Research Development
Consultant



Greet van Meir
Research Administrator



Pat Pokorny
Financial Manager




Maxine Kuroda, MPH, PhD
Epidemiologist / Biostatistician



SITE SUPPORT AND GUIDANCE





Training and Guidance in Practice

- Site training in ICH-GCP in hospital daily practice
- Leading site preparation prior to SIV:
 - Collection essential documents
 - Budget and contract negotiations and finalization
 - Site logistics
 - Trial customized worksheets
- Perform first patient visits in presence of local site staff (on-site support)
- Assist in eCRF completion/query resolution for first patients
- Follow-up on recruitment
- Creation of newsletters for distribution to the sites and client



Unique Advantages

We provide a dedicated team of skilled professionals for each project. Our expertise, experience and flexible structure allows us to anticipate and react to individual client requirements.



Favorable Regulatory Framework and Expertise

The bulk of clinical research is performed in Belgium, known for its favorable regulatory framework. Start-up timelines are among the best in the EU Member States, and allow investigators to activate their sites quickly.



International Collaboration

With several of our USA team members and our location in both USA and Belgium, we are ideally positioned to guide US sponsors into the EU. Due to collaboration with sites in different European countries, we can also provide international Site Management.



Therapeutic Excellence

We are specialized in performing clinical trials in most fields of medicine, with high-end expertise in the fields of Anesthesiology, Critical Care, Pain and Perioperative Medicine.



Tailored Approach

Communication is key to success. Our adaptive structure ensures direct communication lines with the team members that matter. We provide an efficient project team to assist the pharma and devices industry.





Strategic Goals

- Expand our existing research network into a robust research network throughout Belgium and Europe
- Shorten drug development time through a network of research sites and investigators
- Facilitate patient recruitment
- Identify and support additional external sites in starting, managing and executing clinical trials
- Provide a multidisciplinary team for on-site support for potential investigational sites that have the necessary patient pool, but lack the required staff, research experience, or time
- Provide robust regulatory, monitoring, accounting and contracting support to participating research sites



SciMedBE

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