WE CARE

VIVACTIS
SWITZERLAND

CLINICAL STUDY CONCEPTION & AUTHORISATION

Collect credible results
whilst ensuring subject safety

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SWITZERLAND

Who you are:

- A **company** active in the healthcare sector
- A **clinical center/investigator**
- A **health league/NGO…**

… willing to:

- **Conceive a clinical study** that is compliant with ethical and regulatory requirements
- **Prepare the documentation** to obtain the authorisation to run the study

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What we offer you:

CONSULTING

- Medical need identification
- Literature review
- KOL identification
- Study scope definition
- Study outcomes selection
- Clinical trial authorisation procedures

MEDICAL WRITING

- Literature digest
- Study synopsis
- Clinical study authorisation file*: study protocol, informed consent form, questionnaires for participants and HCPs, participants' diaries
- Study report
- Scientific publication & whitepaper

*additional services: study registration on clinicaltrials.gov & electronic submission to the Ethics Committee

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SWITZERLAND

Why us?

- STRONG SCIENTIFIC BACKGROUND
- DEEP KNOWLEDGE OF THE MEDICAL FIELD
- UP-TO-DATE WITH REGULATORY GUIDELINES
- COMPLIANCE WITH REGULATORY AND ETHICAL REQUIREMENTS
- EFFICIENCY
- REACTIVITY
- DEDICATION
- PROFESSIONALISM

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LET'S TALK

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