Are you struggling to find information how to set up a clinical study in Belgium and/or The Netherlands?

Discover and join our training courses!

Despite all efforts for harmonization, setting up a clinical study in Europe remains a challenge. Finding information on the local requirements for regulatory submissions, is scattered and unpractical.

To help you overcome this hurdle, we want to share our years of experience in starting up clinical studies and organize the following theoretical and practical training sessions:

- Regulatory submissions of clinical trials with a medicinal product in Belgium
- Regulatory submissions of clinical investigations with devices in Belgium
- Regulatory submissions of clinical trials with a medicinal product in The Netherlands
- Regulatory submissions of clinical investigations with devices in The Netherlands

The courses guide you through the jumble of local requirements for regulatory submissions to obtain green light to start a study. The courses are illustrated with practical examples and most updated information is guaranteed.

- **Web-based training sessions or Face to Face meetings:**

AML organizes interactive training via web access. Session dates are published on the AML website www.aml-research.be

AML also provides F2F training for companies and institutions on request. Training can be adapted to specific group needs and can be hosted in-company or at AML premises.

For more detailed information, visit our website or contact us at training@aml-research.be or +32 (0)16 78 21 70.