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## **Summary of Qualifications**

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***Professional Experience***

- More than two years experience in clinical trials

***Therapeutic Experience***

- Wound healing (hernia repair, tissue adhesive) Phase IV
- Transplantation (cytomegalovirus, antiviral drugs) Phase III
- Pulmonary embolism (cardiovascular, anticoagulant) Phase III
- Leukemia (philadelphia chromosome, chemotherapy) Phase III
- AML (acute myeloid leukemia, chemotherapy) Phase III
- Perenteral nutrition (vitamines) Phase III
- Low back pain (application study) Phase IV
- Hepatitis C (internal medicine) Phase III
- DNP (diabetic neuropathy, pain drugs) Phase II, Phase III
- PHN (post herpetic neuralgia, antiviral drug) Phase IIa
- Lung cancer (oncology, antibody) Phase IIIa-IIIb
- Inoperable melanoma (oncology, chemo therapy) Phase I-II
- Breast cancer (oncology, antibodies) Phase I, Phase II

## **Professional Experience**

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**Fresenius Biotech GmbH, Munich, Germany**

**Title: Study Manager**

**September 2008 – December 2008**

- Communication between regular authorities, CRO, CRAs and study sites
- Verification and shipment of essential documents
- TMF review
- Review of reports
- Drug import licenses
- Administrative meetings and telephone conferences
- Contract affairs
- Oncology training, 4 days

**INC Research GmbH, Düsseldorf, Germany**

**Title: CRA I**

**September 2007 – September 2008**

- Site selection
- Communication between sponsor and investigator
- IP verification (storage conditions, accountability, etc.)
- Verification of investigator compliance with protocol
- Report of deviations from protocol, GCP, SOPs to investigator and preventing reoccurrence, etc.
- Source documents and CRF verification, verification of data accuracy
- Verification of IC documents and patients/participants enrollment
- Patients/participants safety, AEs/SAEs reporting
- Verification of all essential documents

**Hyperphar Group Germany GmbH, Starnberg, Germany**

**Title: CRA I**

**Dates: 1st October 2006 – 15<sup>th</sup> August 2007**

- Responsible for assisting with the operational administration of clinical research projects
- Feasibilities
- Submission to Ethics Committees and
- Submission to Competent Authorities (BfArM, PEI)
- Performing monitoring visits for wound healing study (hernia repair)
- eCRF training on the job
- SAE- and Data Management training on the job
- GCP-ICH training on the job
- Monitoring training on the job
- Archiving and document tracking (TMF) training on the job
- SOP training certificate

**Hyperphar Group Germany GmbH, Starnberg, Germany**

**Title: CRA-Trainee**

**Dates: 16<sup>th</sup> August 2006– 1<sup>st</sup> October 2006**

**Internship Coromandel Informationssysteme, Germering, Germany**

**Title: Hardware assembler**

**Dates: March 2006-May 2006**

**Title: private school tutor**

**Dates: 2005 - 2006**

**Litec Computer Vertriebs GmbH, Munich, Germany**

**Title: Hardware assembler**

**November 2004 - February 2005**

**Bavarian regional office for fishery, Starnberg, Germany**

**Title: Limnological consultant for the trial „Bio-trout“**

**Dates: 2002**

## **Education**

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**Technical University of Munich (TUM) Munich, Germany**

**Title: Master of Science in Biology (German diploma)**

**Dates: 1993-2002**

**Carl-Spitzweg-Gymnasium Germering, Germering, Germany**

**Diploma from German secondary school qualifying for matriculation**

**Dates: 1984-1993**

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