



## Curriculum Vitae

Personal information **Daniel Bjermo**

### Work experience

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1. Employer: Swedish Medical Products Agency
  - Start date: Oct 2021
  - End date: present
  - Position: Pharmaceutical Inspector GCP
  - Activities: National and international inspections of clinical trials at investigational sites and sponsors/CRO
  - Country: Sweden
2. Employer: LINK Medical Research (formerly PCG Clinical Services)
  - Start date: Mar 2018
  - End date: Oct 2021
  - Position: Quality Assurance Manager
  - Activities: Consultancy including QMS maintenance and revision, conducting GCP audits at CROs, vendors and investigational sites, vendor qualification/re-qualification, quality issue reporting and follow-up, training, and advising on quality-related matters.
  - Country: Sweden
3. Employer: PCG Clinical Services
  - Start date: Sep 2014
  - End date: Mar 2018
  - Position: Manager, Clinical
  - Activities: Leading the team of CRAs in their daily work as monitors in clinical studies. Responsibility for implementation and maintenance of processes and quality documents related to clinical and risk-based monitoring. Working part-time as a Clinical Project Manager, coordinating all activities in full-service clinical and biometrics trials and studies.
  - Country: Sweden
4. Employer: Smerud Medical Research
  - Start date: Sep 2012
  - End date: Sep 2014
  - Position: Clinical Project Manager and Country Manager
  - Activities: Coordination of clinical activities in phase II-III clinical trials and non-interventional studies. Clinical monitoring of phase II-III clinical trials and non-interventional studies. Leading a team of CRAs/monitors, clinical research scientists and biostatisticians.
  - Country: Sweden
5. Employer: Smerud Medical Research
  - Start date: Mar 2012
  - End date: Sep 2012
  - Position: Clinical Project Manager and CRA Manager
  - Activities: Coordination of clinical activities in phase II-III clinical trials and non-interventional studies. Clinical monitoring of phase II-III clinical trials and non-interventional studies. Leading a team of CRAs/monitors. Supervising two students doing their master's degree thesis project in Biomedicine.
  - Country: Sweden
6. Employer: Smerud Medical Research
  - Start date: Aug 2011
  - End date: Mar 2012
  - Position: Senior Clinical Research Associate
  - Activities: Clinical monitoring of phase II-III trials and non-interventional studies
  - Country: Sweden
7. Employer: Quintiles Phase I Services
  - Start date: May 2010
  - End date: Aug 2011
  - Position: Clinical Research Associate
  - Activities: Monitoring of clinical phase I-IIa trials in Sweden
  - Country: Sweden
8. Employer: Orion Pharma
  - Start date: Jan 2008
  - End date: May 2010
  - Position: Clinical Research Associate
  - Activities: Managing and monitoring of clinical phase II-IV trials and non-interventional studies. Supporting investigator-initiated studies
  - Country: Sweden
9. Employer: Orion Pharma
  - Start date: Aug 2007
  - End date: Jan 2008
  - Position: Clinical Trial Assistant
  - Activities: Archiving and administration in clinical phase III-IV trials. Monitoring of clinical phase IV trials
  - Country: Sweden

### Education and training

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1. Subject: Uppsala University

- Start date: Aug 2002
- End date: Jun 2006
- Qualification: MSc. Biomedicine
- Organisation: Uppsala University
- Country: Sweden
- Subjects: E.g. clinical drug development, molecular cell biology, neurology, organic chemistry, immunology, statistics

## Additional information

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Publications

Projects

Memberships

Member of the Swedish Pharmaceutical Society, since 2012

Other Relevant Information