



**Personal Information**

*Damiano Cagnetti*

**Actual Role**

*Study coordinator / Data Entry / Study start-up*

**Education**

Master's degree in Biology (LM-6)

**Previous Professional Experience**

*(se applicabile)*

**Experience in Clinical Research**

*Indicare principali mansioni svolte, una media di quanti studi coordinate/seguite per start-up e in quale fase (I, II e III)*

*Per es. da JD:*

- Knowledge and application of Good Clinical Practice (GCP)
- Knowledge of Standard Operating Procedures (SOP)
- Knowledge of current legislation about Clinical Research
- Data entry activities: compilation of eCRFs and resolution of queries. Use of the main eCRF systems: RDC, Rave EDC, InForm, Veeva, REDCap, OpenClinica ....
- Consultation of reporting systems for the collection of Source Data
- Management of the Investigator Site File/ Investigator Master File
- Planning and management of monitoring visits: from pre-study to close out visit
- Management of IWRS systems

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**Trainings in Clinical Research:**

*Clinical Research Training Course "Missione CRA" (ClinOpsHub , Yghea CRO) 50 hours Clinical Research training course as per MISSIONE CRA the Ministerial Decree 15.11.2011.*