

Investigator Curriculum Vitae

This template may be used by Sponsors of clinical trials as part of the application dossier.

EU CT number: 2024-512278-92

A separate document should be completed and submitted for each site.

This template has been developed and endorsed by the EU Clinical Trials Expert Group to comply with Regulation (EU) No. 536/2014 Clinical Trials on Medicinal Products for Human Use.

Personal Information

Name: PRATI DANIELE
Title: MD
Profession: cardiologist
Current position: Consultant

Professional Registrationⁱ

Registration number: TN 3514
Registration body: Ordine dei Medici della Provincia di Trento
Registration expiry date (if applicable): _____
Registration state/province (if applicable): Italy, Trento

Education and Qualificationsⁱⁱ

Institution name	Qualification	Year
University of Verona	Degree in Medicine and Surgery	2001
University of Verona	Specialization Cardiology disease	2005

Current employment

Institution name: AOUI Verona
Department: Cardiology
Institution address: Piazzale Stefani 1 , 37126 Verona

Investigator Curriculum Vitae

Telephone number: 0039 0458122988

E-mail address: daniele.prati@aovr.veneto.it

Professional experienceⁱⁱⁱ

Position	Institution name and department	Start year	End year
clinical fellow	Unità Operativa di Cardiologia Hospital Policlinico G.B. Rossi , Verona	2005	2007
medical executive	Unità Operativa di Cardiologia Hospital S. Maria del Carmine , Rovereto	2007	2009
medical executive	Unità Operativa di Cardiologia Hospital Policlinico G.B. Rossi , Verona	2009	2012

Relevant clinical trial/study experience^{iv}

Investigator role	Therapeutic area	Type of trial	Year started	Phase	Ongoing End in
Subinvestigator	Acute Coronary Syndrome	Double blind randomized controlled trial	2022	III	2023

Training

Research training (including GCP) ICH GCP - E6	Institution name The global health network	Year obtained
		2022

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Date completed:

22 July 2024

Signature^v:

Dr Daniele Prati



- 1 As per national legislation
- 2 Relevant to be an investigator
- 3 This should cover the preceding 10 years as a maximum
- 4 Idem
- 5 The CTR does not require signing individual documents in the clinical trial application – a request for signature could however be subject to national legislation.