

Investigator Curriculum Vitae

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

EU CT number : **2025-523552-31-00**

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:	Giovanni Adami
Title:	MD, PhD
Profession:	Medical Doctor, Rheumatologist
Current position:	Researcher, Assistant Professor

Professional Registrationⁱ

Registration number:	OMCEO VR08783
Registration body:	Ordine dei medici di Verona
Registration expiry date (if applicable):	NA
Registration state/province (if applicable):	Italy, VR

Education and Qualificationsⁱⁱ

Institution name	Qualification	Year
University of Verona	Medical Doctor (MD)	2015
University of Verona	Rheumatologist	2021
University of Verona	PhD	2023

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Current employment

Institution name: University of Verona

Department: Department of Medicine, Section of Rheumatology

Institution address: Pz Scuro 10, Verona, 37134

Telephone number: +390458123155 - +390458122574

E-mail address: Giovanni.adami@univr.it

Professional experienceⁱⁱⁱ

Position	Institution name and department	Start year	End year
PhD	University of Verona, Department of Medicine, Section of Rheumatology	2018	2019
Post-Doc – Research Fellow	University of Alabama at Birmingham, Division of Clinical Immunology and Rheumatology		
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Relevant clinical trial/study experience^{iv}

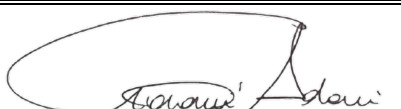
Investigator role	Therapeutic area	Type of trial	Year started	Phase	Ongoing
Sub Investigator	Rheumatology, GIOP (no profit)	Clinical Study	2022	Phase III	No
Sub Investigator	Rheumatology (no profit)	Clinical Study	2023	Phase III	Yes
Sub Investigator	Rheumatology (no profit)	Clinical Study	2022	Phase III	Yes

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Training

Research training (including GCP) Good Clinical Practice (ICH E6 – GCP R3)	Institution name	Year obtained
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Date completed^v: 13/12/2025



ⁱ As per national legislation

ⁱⁱ Relevant to be an investigator

ⁱⁱⁱ This should cover the preceding 10 years as a maximum

^{iv} Idem

^v The CTR does not require signing individual documents in the clinical trial application – a request for signature could however be subject to national legislation.