

Investigator Curriculum Vitae

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

EU CT number : **2022-502678-18-00**

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:	Giacomo Zoppini
Title:	MD
Profession:	Medical doctor
Current position:	Associate professor of Endocrinology

Professional Registrationⁱ

Registration number:	07257
Registration body:	Ordine Medici e Chirurgia di Verona
Registration expiry date (if applicable):	NA
Registration state/province (if applicable):	Verona, Italy

Education and Qualificationsⁱⁱ

Institution name	Qualification	Year
Università degli Studi di Padova	Medical Degree,	1985
Università degli Studi di Modena	PhD: Scienze Endocrinologiche e Metaboliche	1992
Università di Brescia	Italian Board of Endocrinology and Metabolic Diseases	2009

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Current employment

Institution name:	Azienda Ospedaliera Universitaria Integrata Verona
Department:	Endocrinologia, Diabetologia e Malattie del Metabolismo
Institution address:	Piazzale Stefani, 1, Verona, Italia
Telephone number:	390458123115
E-mail address:	Giacomo.zoppini@univr.it

Professional experienceⁱⁱⁱ

Position	Institution name and department	Start year	End year
Associate Professor	Sezione di Endocrinologia, Dipartimento di Medicina, Università di Verona	2014	On going
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Relevant clinical trial/study experience^{iv}

Investigator role	Therapeutic area	Type of trial	Year started	Phase	Ongoing
Sub Investigator	Diabetology	Interventional Trial	2015	Phase III	Yes
Sub Investigator	Diabetology	Interventional Trial	2015	Phase III	No
Sub Investigator	Diabetology	Interventional Trial	2023	Phase II	Yes

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Training

Research training (including GCP)
The Global Health network

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Institution name
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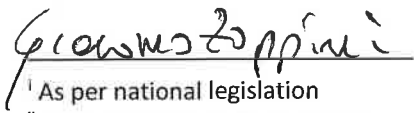
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Year obtained
2023

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Date

completed^v:18/05/2023



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.