

Investigator Curriculum Vitae

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

EU CT number : **2023-504477-21-00**

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:	Emilio Pedrotti
Title:	Prof., MD
Profession:	Medical Doctor
Current position:	Full time University researcher, ophtalmologist and ophtalmic surgeon , Clinica Oculistica dell'Azienda Ospedaliera Integrata di Verona, Policlinico di Borgo Roma (Verona), Italy

Professional Registration¹

Registration number:	n. 0000004754
Registration body:	Albo Provinciale dei Medici Chirurghi di VICENZA (Ordine della Provincia di VICENZA)
Registration expiry date (if applicable):	Click or tap here to enter text.
Registration state/province (if applicable):	Italy/Vicenza

Education and Qualifications²

Institution name	Qualification	Year
Verona University	Specialization in Ophtalmology	2001
Verona University	Degree in "Medicina e Chirurgia"	1997
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Current employment

Institution name: Azienda Ospedaliera Integrata di Verona

Department: Clinica Oculistica

Institution address: Policlinico di Borgo Roma (Verona), Piazzale Ludovico Antonio Scuro, 10 - 37134 Verona (VR), Italy

Telephone number: [REDACTED]

E-mail address: emilio.pedrotti@univr.it

Professional experience³

Position	Institution name and department	Start year	End year
Full time University researcher, ophthalmologist and ophthalmic surgeon	Università di Verona, Clinica Oculistica	2005	-
Professor of pathophysiological optic and squint	Università di Verona	2010	

Relevant clinical trial/study experience⁴

Investigator role	Therapeutic area	Type of trial	Year started	Phase	Ongoing
Choose an item.		Choose an item.		Choose an item.	Choose an item.
Choose an item.		Choose an item.		Choose an item.	Choose an item.
Choose an item.		Choose an item.		Choose an item.	Choose an item.

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<u>Training</u>		
Research training (including GCP)	Institution name	Year obtained
Click or tap here to enter text.	Click or tap here to enter text.	Click or tap here to enter text.
Click or tap here to enter text.	Click or tap here to enter text.	Click or tap here to enter text.

Date completed⁵:

8/06/23

[Signature]

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.