

Investigator Curriculum Vitae

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

EU CT number: **2024-517529-26**

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: **Silvia Romito**

Title: **Dr**

Profession: **Neurologist**

Current position: **MD**

Professional Registrationⁱ

Registration number: **6803**

Registration body: **Medical and Surgeons Register**

Registration expiry date (if applicable): **n.a.**

Registration state/province (if applicable): **Verona-Italy**

Education and Qualificationsⁱⁱ

Institution name	Qualification	Year
University of Verona	Degree in Medicine and Surgery	1998
University of Verona	Specialization in Neurology	2002
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Investigator Curriculum Vitae

Current employment

Institution name: AOUI di Verona
Department: A Neurology
Institution address: P.le Stefani 1 -37126 - Verona
Telephone number: +39 045 8122690
E-mail address: Silvia.romito@aovr.veneto.it

Professional experience^v

Position	Institution name and department	Start year	End year
Assistant	A Neurology Department – AOUI Verona	2002	2006
MD	A Neurology Department – AOUI Verona	2007	Ongoing
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Relevant clinical trial/study experienceⁱⁱⁱ

Investigator role	Therapeutic area	Type of trial	Year started	Phase	Ongoing
SubInv	Amyotrophic Lateral Sclerosis	randomised, multi-centre, double-blind, placebo-controlled, parallel group investigation	2013	II	no
SubInv	Amyotrophic Lateral Sclerosis	randomized, placebo-controlled, double-blind, multi-centre investigation	2010	II	no
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Training

Research training (including GCP)
ICH GCP E6 (R2)

Institution name
The Global Health Network

Year obtained
2025

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Date completed^{iv}: 8th feb 2025

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