

Investigator Curriculum Vitae

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

EU CT number : 2024-510719-31-00

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: Giacomo Allegrini
Title: MD
Profession: Physician
Current position: Oncologist

Professional Registrationⁱ

Registration number: 2404
Registration body: Albo Provinciale dei Medici Chirurghi di LUCCA
Registration expiry date (if applicable): Not applicable
Registration state/province (if applicable): Italy/Lucca

Education and Qualificationsⁱⁱ

Institution name	Qualification	Year
University of Pisa	Specialization in Farmacology	2006
University of Pisa	Specialization in Oncology	1999
University of Pisa	Medicine Degree	1995

Investigator Curriculum Vitae

Current employment

Institution name: Azienda USL Toscana Nord Ovest, Ospedale Civile di Livorno

Department: Department of Oncology

Institution address: Viale Alfieri, 36. 57124-Livorno (LI)

Telephone number: +39-0586223416

E-mail address: giacomo.allegri@uslnordovest.toscana.it

Professional experienceⁱⁱⁱ

Position	Institution name and department	Start year	End year
Oncologist	Department of Oncology, Azienda USL Toscana Nord Ovest, Ospedale Civile di Livorno e VDE-AVC	2018	Ongoing
Oncologist	Ex USL5/Azienda USL Toscana Nord Ovest, Ospedale "F. Lotti" di Pontedera, VDE-AVC	2008	2018

Relevant clinical trial/study experience^{iv}

Investigator role	Therapeutic area	Type of trial	Year started	Phase	Ongoing
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	Oncology		2015		
	Oncology		2014		

Training

Research training (including GCP)
ICH GOOD CLINICAL PRACTICE E6 (R2)

Institution name
Global Health Training Centre

Year obtained
2022

Date completed^v: 17 APR 2024

- 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.