

Investigator Curriculum Vitae

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

EU CT number : **Health-Related Quality of Life in Patients with VEXAS Syndrome: an Italian GIMEMA study**

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: Cinzia Sissa
Title: MD
Profession: Physician
Current position: Hematologist

Professional Registrationⁱ

Registration number: VR7838
Registration body: Ordine dei Medici Chirurghi e Odontoiatri della Provincia di Verona
Registration expiry date (if applicable): N/A
Registration state/province (if applicable): Verona, ITALY

Education and Qualificationsⁱⁱ

Institution name	Qualification	Year
University of Verona, Italy	MD graduation	2007
University of Verona, Italy	Post-graduation in Hematology	2012
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Current employment

Institution name: A.U.O.I. Verona – U.O.C. Ematologia
 Department: DAI Medico Generale
 Institution address: Policlinico G.B.Rossi, P.le L.A.Scuro 10 – 37134 Verona, Italy
 Telephone number: +390458124032
 E-mail address: cinzia.sissa@aovr.veneto.it

Professional experience^{III}

Position	Institution name and department	Start year Click or tap here to enter text.	End year
	ASST Mantova	2012	2023
MD Hematologist		2023	ongoing
MD Hematologist	A.U.O.I. Verona – U.O.C. Ematologia		

Relevant clinical trial/study experience^{IV}

Investigator role	Therapeutic area	Type of trial	Year started	Phase	Ongoing
Sub Investigator	Myelofibrosis	Other research study	2022	Other	No
Sub Investigator	Lymphoma	Clinical Study	2011	Phase III	No
Choose an item.	Click or tap here to enter text.	Choose an item.	Click or tap here to enter text.	Choose an item.	Choose an item.

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Training

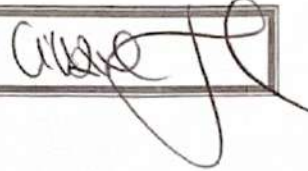
Research training (including GCP)
GCP Certificate

Institution name
Click or tap here to enter text.

Year obtained
2024

Date completed^v:

23.x, 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.