Sub-Investigator Curriculum Vitae

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

EU CT number:

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:	Dr. Alessia Moioli
Title:	Medical degree, Speciality degree in Hematology
Profession:	Hematologist
Current position:	Sub-investigator

Professional Registration[i]

Registration number:	07795
Registration body:	Ordine dei Medici Chirurghi ed Odontoiatri della Provincia di Bergamo
Registration expiry date (if applicable):	NA
Registration state/province (if applicable):	Bergamo, Italy

Education and Qualifications[ii]

Institution name	Qualification	Year
University of Milano	Medicine Degree	2018
University of Verona	Specialistic Degree in Hematology	2025

Sub-Investigator Curriculum Vitae

Current employment

Institution name:	CENTRO RICERCHE CLINICHE DI VERONA SRL
Department:	N/A
Institution address:	P.le L.A. Scuro, 10 - Verona
Telephone number:	[redacted]
E-mail address:	[redacted]

Professional experience[iii]

Position	Institution name and department	Start year	End year
Doctor- Hematologist	Centro Ricerche Cliniche di Verona Srl	Feb 2025	Ongoing
Hematology Fellow	AOUI Verona	Jan 2021	Jan 2025
Fellowship	Cancer Institute of the University College of London for the project *'Molecular and clinical profiling of refractory and early relapsing follicular lymphomas: an unsupervised dual computational approach using co-expression networks and biomarker combinations'* (Prof. T. Marafioti)	Jun 2019	Jul 2019

Relevant clinical trial/study experience[iv]

Sub-Investigator Curriculum Vitae

Sub-Investigator role	Therapeutic area	Type of trial	Year started	Phase	Ongoing
Sub- Investigator	Hematology, Lymphoma	Interventional Trial	2020	Phase II	Yes
Sub-Investigator	Hematology, Lymphoma	Observational Trial	2022	Other	Yes
Sub- Investigator	Hematology, Myeloproliferative neoplasm	Interventional Trial	2022	Phase II	Yes
Sub- Investigator	Hematology, Lymphoma	Interventional Trial	2023	Phase I	Yes
Sub-investigator	Hematology, Myeloproliferative neoplasm	Interventional Trial	2023	Phase I I	Yes
Sub- Investigator	Hematology, Myeloproliferative neoplasm	Interventional Trial	2024	Phase I	Yes
Sub- Investigator	Hematology, Myeloproliferative neoplasm	Interventional Trial	2024	Phase III	Yes
Sub- Investigator	Hematology, Lymphoma	Interventional Trial	2024	Phase I	Yes
Sub- Investigator	Hematology, Lymphoma	Interventional Trial	2024	Phase I	Yes
Sub – Investigator	Hematology, GvHD	Interventional Trial	2025	Phase III	Yes
Sub-investigator	Neurology, Narcolepsy	Interventional Trial	2025	Phase II	Yes
Sub-investigator	Hematology, Lymphoma	Interventional Trial	2025	Phase III	Yes
Sub-investigator	Hematology, Lymphoma	Interventional Trial	2025	Phase I	Yes

Training

Research training (including GCP)	Institution name	Year obtained
ICH GOOD CLINICAL PRACTICE E6 (R3)	Pfizer	2025
ADVANCE LIFE SUPPORT (ALS)	IRC	2022

Sub-Investigator Curriculum Vitae

Date completed[v]:	22-APR-2026

[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.