Investigator Curriculum Vitae

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

EU CT number : 2024-513739-25-00

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:	Andrea Dalbeni
Title:	MD, PhD
Profession:	Physician
Current position:	Consultant at AUOI Verona

Professional Registration[i]

Registration number:	MN2837
Registration body:	Ordine dei Medici e dei Chirurghi
Registration expiry date (if applicable):	
Registration state/province (if applicable):	Italy, MANTOVA

Education and Qualifications[ii]

Institution name	Qualification	Year
Liceo Scientifico Belfiore	High School Diploma	2000
Università degli Studi di Modena e Reggio Emilia	Graduation in Medicine; MD	2006
Università degli Studi di Verona	High graduation	2012

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Università degli Studi di Verona	PhD	2016
Università degli studi di Verona	"Cultore della Materia "(medicina interna)	2019
Università degli studi di Verona	Aggregate professor	2022
MIUR	Professor (Abilitazione Professore di II fascia)	2023

Current employment

Institution name:	AOUI Verona
Department:	UOC Medicina Generale C
Institution address:	p.le scuro 10, 37134 Verona
Telephone number:	+390458124414
E-mail address:	andrea.dalbeni@aovr.veneto.it

Professional experience[iii]

Position	Institution name and department	Start year	End year
Consultant	AOUI Verona- Medicine	2015	On going
High speciality	AOUI VERONA	2025	On going

Relevant clinical trial/study experience[iv]

Investigator role	Therapeutic area	Type of trial	Year started	Phase	Ongoing
Sub Investigator	HCC (no profit)	Clinical Study	2024	Other	Yes
Principal Investigator	Heart Failure (no profit)	Clinical Study	2011	Phase III	No

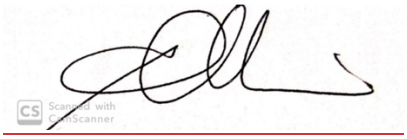
Investigator Curriculum Vitae

Principal Investigator	HCC (no profit)	Clinical Study	2024	Other	Yes

Training

Research training (including GCP)	Institution name	Year obtained
PhD	Università degli Studi di Verona	2016
GCP (E6R3)	Syneos Health	2025

Date completed[v]: 16/12/2025



[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.