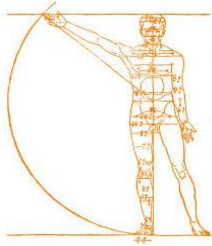


CURRICULUM VITAE – *Stefano Milleri*

<b>CURRICULUM VITAE</b>
<b>STEFANO MILLERI</b>

<b>GENERAL INFORMATION</b>	
Date of birth	07 December 1958
Place of birth	Citta' di Castello (PG)
Nationality	Italian
Mother tongue Language	Italian
Present Job Position	Medical Scientific Director Direttore Sanitario Head of Operation Unit (ad interim) At Centro Ricerche Cliniche di Verona s.r.l., P.le L.A.Scuro 10, 37134 Verona

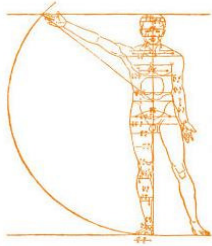
<b>EDUCATION</b>	
Education	PhD in Neuroscience
Institution – City (Country)	Padua University, Faculty of Medicine
Date	2008
Education	Specialization in Geriatrics
Institution – City (Country)	Catholic University in Rome, Faculty of Medicine
Date	1993
Education	Specialization in Lung Diseases
Institution – City (Country)	Catholic University in Rome, Faculty of Medicine
Date	1989
Education	Degree in Medicine
Institution – City (Country)	Catholic University in Rome, Faculty of Medicine
Date	1985



## CURRICULUM VITAE – *Stefano Milleri*

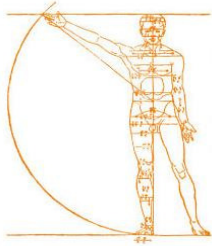
<b>LANGUAGE SKILLS</b>			
Language	English	Level	3
<p>Level 1: Beginner understands read and spoken language, not able to clearly formulate his thoughts in the given language, makes grammar mistakes and uses very basic vocabulary when speaking or writing.</p> <p>Level 2: Speaking skills are better but the vocabulary used and sentence structures are still basic; makes pronunciation mistakes, lacks fluency.</p> <p>Level 3: Comfortable speaker and writer, can build more complex structures and formulate thoughts in a clear manner, makes some grammar mistakes.</p> <p>Level 4: Experienced and fluent user, pronunciation &amp; accent are correct, speaking rhythm is regular, writing skills are developed, language used is rich.</p>			

<b>PROFESSIONAL EXPERIENCE/CAREER HISTORY</b>	
Company	Centro Ricerche Cliniche di Verona s.r.l.
Job Position	Medical Scientific Director and Direttore Sanitario
Date (from-to)	Dec 2019
Main tasks and Responsibilities	<ul style="list-style-type: none"> <li>Providing medical and scientific leadership to the research activities of the unit</li> <li>Ensuring that all Unit operations were covered by Method Sheets or Standard Operation Procedures according to the GCP requirements</li> <li>Ensuring the allocation of qualified physicians (using internal and external staff), appropriately trained, as principal investigators or co-investigators to carry out all unit clinical activities</li> <li>Ensuring the full integration of the CRC with the Hospital Units</li> <li>Principal Investigator or Sub-Investigators in more than 100 Phase 1 and 2 studies in different therapeutic areas performed according to GCP requirements.</li> <li>Co-author of dossiers submitted to European Regulatory Agencies and to FDA</li> <li>Responsible for Clinical Risk Management</li> <li>Guaranteeing the CRC compliance with Italian law related to medical activities (DPR 27/03/1969)</li> </ul>
Company	Centro Ricerche Cliniche
Job Position	Head of Operation Unit (ad interim)
Date (from-to)	May 2016
Main tasks and Responsibilities	<ul style="list-style-type: none"> <li>Supervision of project management and regulatory activities</li> <li>Ensuring all unit activities are performed according to local, ethical and regulatory standards</li> </ul>
Company	Centro Ricerche Cliniche
Job Position	Medical Scientific Director
Date (from-to)	June 2016 – Dec 2019
Main tasks and Responsibilities, if relevant for your present position:	<ul style="list-style-type: none"> <li>Providing medical and scientific leadership to the research activities of the unit</li> <li>Ensuring that all Unit operations were covered by Method Sheets or Standard Operation Procedures according to the GCP requirements</li> </ul>



## CURRICULUM VITAE – *Stefano Milleri*

	<ul style="list-style-type: none"> <li>• Ensuring the allocation of qualified physicians (using internal and external staff), appropriately trained, as principal investigators or co-investigators to carry out all unit clinical activities</li> <li>• Ensuring the full integration of the CRC with the Hospital Units</li> <li>• Principal Investigator or Sub-Investigators in more than 100 Phase 1 and 2 studies in different therapeutic areas performed according to GCP requirements.</li> <li>• Co-author of dossiers submitted to European Regulatory Agencies and to FDA.</li> </ul>
Company	Centro Ricerche Cliniche
Job Position	Scientific Director
Date (from-to)	Jan 2006 - Jun 2016
Main tasks and Responsibilities, if relevant for your present position:	<ul style="list-style-type: none"> <li>• Providing medical and scientific leadership to the research activities of the unit</li> <li>• Training and developing therapeutically medical staff for the supervision and carrying out clinical studies</li> <li>• Ensuring that all Unit operations were performed according to the GCP requirements</li> <li>• Ensuring the allocation of qualified physicians (using internal and external staff), appropriately trained, as principal investigators or co-investigators to carry out all unit clinical activities</li> </ul>
Company	GlaxoSmithKline (former Glaxo, GlaxoWellcome)
Job Position	Director of Clinical Pharmacology & Experimental Medicine Unit
Date (from-to)	2002 - 2005
Main tasks and Responsibilities, if relevant for your present position:	<ul style="list-style-type: none"> <li>• Providing medical and scientific leadership to the research activities of the unit and the development of specialist technologies and humane disease models to meet needs of Discovery Medicine</li> <li>• Training and developing therapeutically medical staff for the supervision and carrying out clinical studies</li> <li>• Ensuring that all Unit operations were covered by Method Sheets or Standard Operation Procedures according to the GCP requirements</li> <li>• Ensuring the allocation of qualified physicians (using internal and external staff), appropriately trained, as principal investigators or co-investigators to carry out all unit clinical activities</li> <li>• Developing new standards of practices for the application of novel technologies and humane disease models</li> <li>• Contributing to the Psychiatric Leadership Team.</li> </ul>
Company	GlaxoSmithKline (former Glaxo, GlaxoWellcome)
Job Position	Physician Responsible for Clinical Pharmacology Unit
Date (from-to)	1992 - 2002
Main tasks and Responsibilities, if relevant for your present position:	<ul style="list-style-type: none"> <li>• Ensuring the allocation of qualified physicians (using internal and external staff), appropriately trained, as principal investigators or co-investigators to carry out all unit clinical activities</li> </ul>



## CURRICULUM VITAE – *Stefano Milleri*

	<ul style="list-style-type: none"> <li>• Contributin to protocol definition</li> <li>• Safety data revision</li> </ul>
Company	Internal Medicine Department at Catholic University Hospital in Rome.
Job Position	Attending physician at the Respiratory Physiopathology Unit
Date (from-to)	1985 - 1991
Main tasks and Responsibilities, if relevant for your present position:	Responsible for outpatient activities with Asthmatic and COPD subjects Sub Investigator in several clinical trials

### **PUBLICATIONS**

Co-author of more than 34 full papers published in peer review journal and more than 43 abstracts

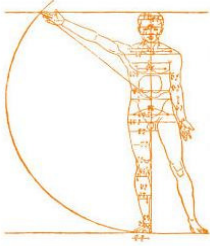
### **OTHER CERTIFICATIONS RELEVANT FOR YOUR POSITION/ ORGANIZATIONS MEMBERSHIP**

Intermediate Life Support (ILS) (Italian Resuscitation Council)

Advanced Life Support (ALS) course (Italian Resuscitation Council): October 2019

### **OTHER INFORMATION RELEVANT FOR YOUR POSITION**

- Acting as Principal Investigator in more than 70 clinical studies (with patients mainly Respiratory, Vaccines and with Healthy volunteers)
- Acting as Sub-investigator in more than 100 clinical studies (mainly Neurology, Oncology, Hematology, CNS)
- Visiting professor at training course of Clinical Pharmacology and Tossicology, Florence University from 2012 up to now
- Lecturer, Master in “Ricerca e Sviluppo Pre-clinico e Clinico dei Farmaci” organized by University of Milan Bicocca from 2009 up to 2020
- Lecturer at Master in “Sperimentazione Clinica dei Farmaci in Medicina Internistica, Ematologia e Oncologia”, organised by Department of Pharmacology, Pisa University from 2004 up to now
- Lecturer, Master in “Regulatory compliance”, organised by Department of Pharmacology, Catania University 2004-2005 and 2005-2006 academic years
- Lecturer at “Indirizzo di Eccellenza in Biomedicina” organized by Istituto di Studi Superiori dell’Università di Genova, in 2015 and 2016
- Lecturer, Master in “Regulatory compliance”, organised by Department of Pharmacology, Catania University 2004-2005 and 2005-2006 academic years



## CURRICULUM VITAE – *Stefano Milleri*

- Lecturer, Master in “farmacovigilanza, farmacoepidemiologia, farmacoconomia e real world evidence”, organized by Verona University, from 2022
- Registered to the Ordine dei Medici di Verona n. 7682 from 2006
- Member of Italian Society of Pharmacology from 2007

*I authorize the treatment of my data according to the applicable local regulation about data protection and any further amendments*