

PERSONAL INFORMATION

Claudio Governatori



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Sex: M | Date of birth: 03/09/1958 | Nationality: Italian

WORK EXPERIENCE

From April 2005 to present: Independent consultant – Holder studio Governatori

In January 2005 I started a business as consultant for pharmaceutical companies: this activity focuses on development /updating of dossier and on Regulatory Affairs services.

Sector: Pharmaceutical consulting

From March 2002 to April 2005: Technology transfer manager at Pulitzer Italiana - Rome

Strategic regulatory guidance for technology transfer concerning the movement of production activities to other manufacturing sites.

Sector: Pharmaceutical

From Sept. 1991 to March 2002: Production manager at Pulitzer Italiana - Rome

Direct pharmaceutical industry management experience and leadership in Manufacturing and Development of drug products.

Sector: Pharmaceutical

From Sept. 1987 to July 1991: Project Leader at Pharmaceutical Technology Department of Chiesi Farmaceutici - Parma

Responsible of Pharmaceutical Technology for the liquid forms, semisolids and parenterals.

In this position I attended to pharmaceutical development of products, from formulation to industrial scale up.

Sector: Pharmaceutical

From April. 1984 to July 1987: Assistant to the Plant manager at Biomedica Foscama S.p.A.-
Ferentino (FR)

Assistant to the Plant manager concerning the production aspects.

Sector: Pharmaceutical

EDUCATION

September 17th 1982: Chemistry Master degree
University of Modena - Italy

July 1977: Scientific high school degree
Montefiascone (VT) - Italy

MAIN TRAINING

September 1987: 1st Master course Biopharmaceutics and Pharmacokinetics
University of Pavia

September 2003: Drug Master File
Istituto Internazionale di Ricerca - Milano

March 2005: Dissolution Rate: Quality and Bioequivalence Aspects
Parenteral Drug Association - Roma

November 2006: Product Quality Review
Parenteral Drug Association - Roma

February 2008: Good Practises of Pharmacovigilance
Pharma Education Center – Pomezia (RM)

July 2008: Module 3 and Quality variations: regulatory compliance
Temas - Milano

April 2010: Pharmacovigilance: Communitarian Regulation updates
Temas - Milano

- March 2011: User training course Eudravigilance and electronic transmission of ICSRs
DIA – S. Marino
- November 2011: Extended Eudravigilance Medicinal Product Dictionary (XEVMPPD)
DIA – S. Marino
- Dec. 2012 – Oct. 2013: Regulatory Affairs Master
Agenzia Italiana del Farmaco / Pharmacology Department of “La Sapienza” University – Roma

PERSONAL SKILLS

Mother tongue : Italian

Other language(s)	UNDERSTANDING		SPEAKING		WRITING
	Listening	Reading	Spoken interaction	Spoken production	
English	B1	B2	B1	B2	C1

ADDITIONAL INFORMATION

QP qualification: “Qualified person” status recognized by Italian Health Authority in January 2003 by act N. IDT – 7/2003, according to the Italian Law N. 178 of May 29th 1991 as amended

- Associations :
- Italian Professional Association of chemists (L.U.A.M.)
 - AFI - Associazione Farmaceutici Industria
 - ANCTF - Associazione nazionale Tecnologi Farmaceutici
 - AICF - Associazione Italiana Consulenti del farmaco.