

PERSONAL INFORMATION

First name(s) / Surname(s)	Eliseo Di Giosaffatte
Address(es)	Siena, Italy
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E-mail	eliseo.digosaffatte@gmail.com
Nationality	Italian
Date of birth	1961
Gender	Male

WORK EXPERIENCE

Dates	From Jan, 1st 2018 to 30 Sep. 2026
Occupation or position held	Global Supplier Quality Auditor
Main activities and responsibilities	I was performing supplier audits for the whole GSK group
Name and address of employer	GSK Vaccines, Supplier Quality Audit & Compliance
Type of business or sector	Biopharmaceutical company, Vaccines producer

Dates	From 01 March 2015 to 31 December 2017
Occupation or position held	Supplier Quality Unit Manager
Main activities and responsibilities	I'm responsible for the whole supplier qualification process; my group has to inspect and to monitor 98 supplier, 3000 different materials We are also deeply involved in all the quality issues arisen from supplier quality issues. Many routine and for cause audits performed each year; around 40 audits, as average each year, for the whole team. My group is made by 4 employees plus me
Name and address of employer	GSK Vaccines s.r.l., Rosia, Italy
Type of business or sector	Biopharmaceutical company, Vaccines producer

Dates	From 01 May 2011 to March 2015
Occupation or position held	Supplier Quality Unit Manager
Main activities and responsibilities	I was responsible for the whole supplier qualification process; my group had to inspect and to monitor 98 supplier, 3000 different materials We were also deeply involved in all the quality issues arisen from supplier quality issues Many routine and for cause audits performed each year; around 40 audits, as average each year, for the whole team. My group was made by 4 employees plus me
Name and address of employer	Novartis Vaccines & Diagnostic, Rosia, Italy
Type of business or sector	Biopharmaceutical company, Vaccines producer
Dates	From October 2006 to April 2011
Occupation or position held	Head of Compliance , Italy
Main activities and responsibilities	I was responsible of the all supplier qualification process plus host all the regulatory/Divisional inspections the site received. In two year I managed the preparation , the execution the observations' response and Follow up of 46 regulatory inspections, successfully 4 FDA inspections, several Canadian MoH, Italian MoH, HALAL certification inspections, Brazil, Mexico, EMA, GCC, inspector were received 10 people were with me
Name and address of employer	Novartis Vaccines & Diagnostic, Rosia, Italy
Type of business or sector	Biopharmaceutical company, Vaccines producer
Dates	From October 2002 to September 2006
Occupation or position held	Head of Corporate regulatory Compliance, Eu & ROW
Main activities and responsibilities	I was responsible for the Compliance activities of the Chiron sites in Europe and in the "rest of world" sites. I had collaborators (in total 15) in Italy, Germany and UK. I was responsible for Compliance in the vaccines BU, in the Blood testing one and also in the Biopharmaceutical BU I was in charge with internal audit, External audits, Hosting of regulatory bodies inspections and in GMP training. Compliance assessor for new projects
Name and address of employer	Chiron Corporation
Type of business or sector	Biopharmaceutical, Blood Testing devices and Vaccines producer multinational company
Dates	From June 1997 to September 2002
Occupation or position held	TQM & Compliance Manager
Main activities and responsibilities	I was responsible for internal and external auditing , plus the compilation of the Site Master file
Name and address of employer	Chiron S.p.A. Siena, Italy
Type of business or sector	Biopharmaceutical company, Vaccines producer

Dates	From Feb 1987 to June 1997
Occupation or position held	Head of Chemical lab (first two years) Head of analytical method validation (1 year) Head of liquid formulation production unit (two years) Head of industrial engineering (one year) Quality assurance manager (Last 4 years)
Main activities and responsibilities	During the last four years I was called to build a completely new QA department. In that period Janssen received successfully his first PAI for a liquid formulation
Name and address of employer	Janssen- Cilag, Latina, Italy
Type of business or sector	Pharmaceutical company, Tablets, cream and syrups producer

SKILLS AND COMPETENCES

Personal skills and competences	Degree on CTF (Pharmaceutical chemistry) 110/110
Mother tongue(s)	Italian
Other language(s)	Fluent English
Social skills and competences	Easy going people, innovative and creative.
Organisational skills and competences	Very good team builder, motivated and able to motive others. Flexible and "Open mind" person
Technical skills and competences	Excellent knowledge in GMP regulations, in aseptic and in non-aseptic operation, in Biotech and in Pharma companies. Excellent knowledge of Sterile primary glass containers production process& quality matters, Increasing knowledge in Medical Device Regulation & Auditing.
Driving licence	B Licence
Additional information	J&J certified auditor Novartis Certified auditor, Gold status for GMP audits GSK certified auditor More than 400 inspections performed around the world. Audits covered Pharma companies, development centres and all different type of suppliers and services providers.